

## SEARCH REQUEST FORM

1.737

Requestor's  
Name:

PHILLIP GAMBEL

Serial

Number:

08/487283

Date:

1/28/97

Phone:

303.3997

Art Unit:

1806

## Search Topic:

Please write a detailed statement of search topic. Describe specifically as possible the subject matter to be searched. Define any terms that may have a special meaning. Give examples or relevant citations, authors, keywords, etc., if known. For sequences, please attach a copy of the sequence. You may include a copy of the broadest and/or most relevant claim(s).

SEQ / INTERFERENCE SEARCH

SEQ ID NO. 1 + 2  
= =

REQUEST IN

Three

## STAFF USE ONLY

Date completed:

1/28/98

Searcher:

m

Terminal time:

10

Elapsed time:

CPU time:

Total time:

10

Number of Searches:

1

Number of Databases:

5

## Search Site

☐ STIC☒ CM-1☐ Pre-S

## Type of Search

☐ N.A. Sequence☒ A.A. Sequence <sup>x2</sup>☐ Structure☐ Bibliographic

## Vendors

☒ IG m☐ STN☐ Dialog☐ APS☐ Geninfo☐ SDC☐ DARC/Questel☐ Other



```
#cross-references MUID:88209511
#accession A27689
#molecule_type mRNA
##residues 412-1676 ##label WET
##cross-references GB:M18879
REFERENCE
A01267
#authors Fernandez, H.N.; Hugli, T.E.
#journal J. Biol. Chem. (1978) 253:6955-6964
#title Primary structural analysis of the polypeptide portion of
human C5a anaphylatoxin. Polypeptide sequence determination
and assignment of the oligosaccharide attachment site in
C5a.
#cross-references MUID:79005687
#accession A01267
#molecule_type protein
##residues 678-751 ##label FER
REFERENCE
A01266
#authors Lundwall, A.B.; Wetzel, R.A.; Kristensen, T.; Whitehead,
A.S.; Woods, D.E.; Ogden, R.C.; Colten, H.R.; Tack, B.F.
#journal J. Biol. Chem. (1985) 260:2108-2112
#title Isolation and sequence analysis of a cDNA clone encoding the
fifth complement component.
#cross-references MUID:85130937
#accession A01266
#molecule_type mRNA
##residues 412-854,
'SIALSPRLPCNGKTSCHCKLRPGSDSPASQVAGTGTTHHAQPT'
##label LUN
##cross-references GB:K02874
##note the carboxyl-terminal part of the sequence in this
report appears to be derived from translation of an
AUI repeat sequence
REFERENCE
S15121
#authors Bohnsack, J.F.; Mollison, K.W.; Buko, A.M.; Ashworth, J.C.;
Hill, H.R.
#journal Biochem. J. (1991) 273:635-640
#title Group B streptococci inactivate complement component C5a by
enzymic cleavage at the C-terminus.
#cross-references MUID:91144547
#contents annotation
#COMMENT Complement C5 contains two disulfide-linked chains, formed by
removal of four basic residues. C5 convertase releases C5a
anaphylatoxin from the amino end of the alpha chain, generating
C5b (beta and alpha' chains).
#COMMENT Activation of C5 initiates the spontaneous assembly of the late
complement components, C5-C9, into the membrane attack complex.
C5b has a transient binding site for C6. The C5b-C6 complex is
the foundation upon which the membrane attack complex is
assembled.
#COMMENT C5a has potent spasmogenic and chemotactic activity.
GENETICS
#gene GDB:C5
##cross-references GDB:119734
#map_position 9q33-9q33
CLASSIFICATION #superfamily alpha-2-macroglobulin
#complement alternate pathway; complement pathway; cytolysis;
glycoprotein; inflammation; membrane attack complex; plasma
FEATURE
1-18 #domain signal sequence #status predicted #label SIC\
19-673, 678-1676 #product complement C5 #status predicted #label MAT\
19-673, 752-1676 #product C5b #status predicted #label C5B\
19-673 #product complement C5 and C5b beta chain #status
predicted #label C5B8\
678-1676 #product complement C5 alpha chain #status predicted
```

```
#label C5A\
#product C5a anaphylatoxin #status experimental #label
C5A\
#product C5b alpha' chain #status predicted #label C5BA\
678-751
752-1676
567-810, 634-669,
698-724, 699-731,
711-732, 866-1527,
1101-1159,
1375-1505,
1405-1474,
1520-1525,
1532-1606,
1553-1676,
1654-1657
741
#disulfide bonds #status predicted\
#binding site carbohydrate (Asn) (covalent) #status
experimental\
#cleavage site Arg-Leu (C5 convertase) #status
experimental\
#binding site carbohydrate (Asn) (covalent) #status
predicted
SUMMARY #length 1676 #molecular-weight 188330 #checksum 3858
Query Match 100.0%; Score 141; DB 2; Length 1676;
Best Local Similarity 100.0%; Pred. No. 3,46e-20;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db 872 vidhgtkscvcrqkvsgs 892
|||||
Qy 1 VDRQGTGKSKVRQKVEGSS 21
|||||
RESULT 2
ENTRY #type complete
TITLE complement C5 precursor - mouse
CONTAINS C5a anaphylatoxin; C5b
ORGANISM #formal_name Mus musculus #common_name house mouse
DATE 19-Nov-1988 #sequence_revision 15-Oct-1994 #text_change
16-Feb-1997
ACCESSIONS A35530; A27538; A40429
REFERENCE A35530
#authors Wetzel, R.A.; Fleischer, D.T.; Haviland, D.L.
#journal J. Biol. Chem. (1990) 265:2435-2440
#title Deficiency of the murine fifth complement component (C5). A
2-base pair gene deletion in a 5'-exon.
#cross-references MUID:90153853
#accession A35530
#molecule_type mRNA
##residues 1-215, 'L' ##label WET
##cross-references GB:J05234
REFERENCE A27538
#authors Wetzel, R.A.; Ogata, R.T.; Tack, B.F.
#journal Biochemistry (1987) 26:737-743
#title Primary structure of the fifth component of murine
complement.
#cross-references MUID:87185363
#accession A27538
#molecule_type mRNA
##residues 'PGL', 44-1680 ##label WET2
REFERENCE A40429
#authors Haviland, D.L.; Haviland, J.C.; Fleischer, D.T.; Wetzel, R.A.
#journal J. Biol. Chem. (1991) 266:11818-11825
#title Structure of the murine fifth complement component (C5) gene.
A large, highly interrupted gene with a variant donor
splice site and organizational homology with the third and
```

Jan 28 12:17

US-08-487-283A-1.rpr

5

```
#cross-references MUID:91268053
#accession R40429
#molecule_type DNA
#residues 1-15 ##label HAV
##cross-references GB:M64852
COMMENT Complement C5 contains two disulfide-linked chains, formed by
removal of four basic residues. C5 convertase releases C5a
anaphylatoxin from the amino end of the alpha chain, generating
C5b (beta and alpha' chains).
COMMENT Activation of C5 initiates the spontaneous assembly of the late
complement components, C5-C9, into the membrane attack complex.
C5b has a transient binding site for C6. The C5b-C6 complex is
the foundation upon which the membrane attack complex is
assembled.
COMMENT C5a has potent spasmogenic and chemotactic activity.
GENETICS
#map_position 2
#introns 22/3; 86/3; 140/3; 164/3; 195/2; 223/1; 253/2; 291/3; 334/1;
372/3; 434/3; 502/3; 572/3; 622/3; 667/1; 691/1; 757/1;
787/2; 812/1; 858/3; 934/3; 955/1; 985/1; 1056/1; 1081/2;
1134/3; 1166/3; 1224/1; 1292/3; 1343/3; 1364/3; 1392/1;
1411/2; 1445/3; 1470/3; 1506/1; 1534/1; 1564/1; 1592/1;
1637/2
CLASSIFICATION #superfamily alpha-2-macroglobulin
KEYWORDS complement alternate pathway; complement pathway; cytolysis;
glycoprotein; inflammation; membrane attack complex; plasma
FEATURE
1-18 #domain signal sequence #status predicted #label SIG\
19-674,679-1679 #product complement C5 #status predicted #label MAT\
19-674,756-1679 #product C5b #status predicted #label C5B\
19-674 #product complement C5 and C5b beta chain #status
predicted #label C5BB\
#product complement C5 alpha chain #status predicted
#label C5A\
679-755 #product C5a anaphylatoxin #status predicted #label C5T\
#product C5b alpha' chain #status predicted #label C5BA\
756-1679
567-814,635-670,
702-728,703-735,
715-736,870-1531,
1105-1163,
1379-1509,
1409-1478,
1524-1529,
1536-1609,
1557-1679,
1657-1660
#disulfide bonds #status predicted\
#binding_site carbohydrate (Asn) (covalent) #status
predicted
SUMMARY #length 1680 #molecular-weight 188876 #checksum 3888
Query Match 48.9%; Score 69; DB 2; Length 1680;
Best Local Similarity 47.1%; Pred. No. 9.17e-03;
Matches 8; Conservative 7; Mismatches 2; Indels 0; Gaps 0;
Db 880 htsrparcvfgriegss 896
: ::::|::|::|::|
Qy 5 QGTRSSKVRQKVEGSS 21
RESULT 3
ENTRY #type complete
TITLE hypothetical protein - Synecchocystis sp. (PCC 6803)
ORGANISM #formal_name Synecchocystis sp.
```

Jan 28 12:17

US-08-487-283A-1.rpr

6

```
#variety PCC 6803
DATE 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change
25-Apr-1997
ACCESSIONS S76070
REFERENCE S74322
#authors Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.;
Nakamura, Y.; Miyajima, N.; Hiroseawa, M.; Sugitara, M.;
Sasamoto, S.; Kimura, T.; Hosouchi, T.; Matsuno, A.;
Muraki, A.; Nakazaki, N.; Naruo, K.; Okumura, S.; Shimpou,
S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.;
Yasuda, M.; Tabata, S.
#journal DNA Res. (1996) 3:109-136
#title Sequence analysis of the genome of the unicellular
cyanobacterium Synecchocystis sp. PCC6803. II. Sequence
determination of the entire genome and assignment of
potential protein-coding regions.
#accession S76070
#status preliminary
#molecule_type DNA
#residues 1-213 ##label KAN
##cross-references EMBL:D63999
#note the nucleotide sequence was submitted to the EMBL Data
Library, June 1996
SUMMARY #length 213 #molecular-weight 22745 #checksum 2191
Query Match 44.0%; Score 62; DB 11; Length 213;
Best Local Similarity 35.0%; Pred. No. 2.43e-01;
Matches 7; Conservative 7; Mismatches 6; Indels 0; Gaps 0;
Db 92 ilayigakascdikpkvss 111
: :|::|::|::|
Qy 1 VIDHQGTRKSKVRQVEGS 20
RESULT 4
ENTRY #type complete
TITLE lactoferrin binding protein - Neisseria meningitidis
ORGANISM #formal_name Neisseria meningitidis
DATE 16-Feb-1995 #sequence_revision 12-May-1995 #text_change
12-May-1995
ACCESSIONS S49087
REFERENCE S49087
#authors Pettersson, A.M.; Klarenbeek, X.Y.Z.; van Deurzen, X.Y.Z.;
Poolman, X.Y.Z.; Tomassen, X.Y.Z.
#submission submitted to the EMBL Data Library, June 1994
#description Molecular characterization of the structural gene for the
lacto-ferrin receptor of the meningococcal strain H44/76.
#accession S49087
#status preliminary
#molecule_type DNA
#residues 1-940 ##label PET
##cross-references EMBL:X79838
SUMMARY #length 940 #molecular-weight 105347 #checksum 8194
Query Match 41.8%; Score 59; DB 10; Length 940;
Best Local Similarity 50.0%; Pred. No. 9.30e-01;
Matches 7; Conservative 5; Mismatches 2; Indels 0; Gaps 0;
Db 592 rsrkcvprkingen 605
: |::|::|::|
Qy 8 KSSKVRQKVEGSS 21
RESULT 5
```



Jan 28 12:17

US-08-487-283A-1.rpf

7

```

ENTRY      A56678      #type complete
TITLE      yemanuclein-alpha - fruit fly (Drosophila melanogaster)
ORGANISM   #formal name Drosophila melanogaster
DATE       08-Jul-1995 #sequence_revision 03-Aug-1995 #text_change
          16-Feb-1997
ACCESSIONS A56678; S22146
REFERENCE  #authors    Alt-Ahmed, O.; Bellon, B.; Capri, M.; Joblet, C.;
          Thomas-Delaage, M.
#journal   Mech. Dev. (1992) 37:69-80
#title     The yemanuclein-alpha: a new Drosophila DNA binding protein
          specific for the oocyte nucleus.
#accession A56678
#status    preliminary
#molecule_type DNA
#residues  1-1002 #label A1E
#cross-references GB:X63503
GENETICS
#gene      FlyBase:yem4agr
#cross-references FlyBase:FBgn0005596
#introns   80/3; 154/3; 428/1 477/2; 557/2
KEYWORDS   DNA binding; oocyte
SUMMARY    #length 1002 #molecular-weight 109310 #checksum 4278

Query Match 41.1%; Score 59; DB 15; Length 1002;
Best Local Similarity 50.0%; Pred. No. 9.30e-01;
Matches 6; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

Db 47 tktakirikid 58
   :::| | | |
Qy 7 TKSSKCVQKVE 18

RESULT 6
ENTRY 149364 #type complete
TITLE protein tyrosine phosphatase - mouse
ORGANISM #formal name Mus musculus #common name house mouse
DATE 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change
          02-Jul-1996
ACCESSIONS 149364
REFERENCE 149364
#authors  Wishart, M.J.; Denu, J.M.; Williams, J.A.; Dixon, J.E.
#journal  J. Biol. Chem. (1995) 270:26782-26785
#title    A single mutation converts a novel-phosphotyrosine binding
          domain into a dual-specificity phosphatase.
#accession 149364
#status    preliminary; translated from GB/EMBL/DBDJ
#molecule_type mRNA
#residues  1-205 #label RES
#cross-references EMBL:U34973; NID:g1063624; CDS PID:g1063625
SUMMARY    #length 205 #molecular-weight 23683 #checksum 2745

Query Match 41.1%; Score 58; DB 16; Length 205;
Best Local Similarity 25.0%; Pred. No. 1.44e+00;
Matches 5; Conservative 9; Mismatches 6; Indels 0; Gaps 0;

Db 51 ilqthgithicirqnean 70
   :::| | | | | | | |
Qy 1 VIDHQGTSKSCVRQKVE 20

RESULT 7
ENTRY 149365 #type complete
TITLE protein tyrosine phosphatase - mouse

```

Jan 28 12:17

US-08-487-283A-1.rpf

8

```

ORGANISM   #formal name Mus musculus #common name house mouse
DATE       02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change
          02-Jul-1996
ACCESSIONS 149365
REFERENCE 149364
#authors  Wishart, M.J.; Denu, J.M.; Williams, J.A.; Dixon, J.E.
#journal  J. Biol. Chem. (1995) 270:26782-26785
#title    A single mutation converts a novel-phosphotyrosine binding
          domain into a dual-specificity phosphatase.
#accession 149365
#status    preliminary; translated from GB/EMBL/DBDJ
#molecule_type mRNA
#residues  1-223 #label RES
#cross-references EMBL:U34973; NID:g1063624; CDS PID:g1063626
GENETICS
#introns   168/3
SUMMARY    #length 223 #molecular-weight 25416 #checksum 359

Query Match 41.1%; Score 58; DB 16; Length 223;
Best Local Similarity 25.0%; Pred. No. 1.44e+00;
Matches 5; Conservative 9; Mismatches 6; Indels 0; Gaps 0;

Db 51 ilqthgithicirqnean 70
   :::| | | | | | | |
Qy 1 VIDHQGTSKSCVRQKVE 20

RESULT 8
ENTRY 168524 #type complete
TITLE ribosomal protein L34 - human
ORGANISM #formal name Homo sapiens #common name man
DATE 24-May-1996 #sequence_revision 24-May-1996 #text_change
          16-Feb-1997
ACCESSIONS 168524
REFERENCE 154209
#authors  Rommens, J.M.; Durocher, F.; McArthur, J.; Tonin, P.;
          Leblanc, J.
#journal  Genomics (1995) 28:530-542
#title    Generation of a transcription map at the HSD17B locus
          centromeric to BRCA1 at 17q21.
#accession 168524
#status    preliminary; translated from GB/EMBL/DBDJ
#molecule_type mRNA
#residues  1-117 #label RES
#cross-references GB:L38941; NID:g1008855; CDS PID:g1008856
CLASSIFICATION #superfamily rat ribosomal protein L34
SUMMARY    #length 117 #molecular-weight 13305 #checksum 4392

Query Match 40.4%; Score 57; DB 8; Length 117;
Best Local Similarity 41.7%; Pred. No. 2.22e+00;
Matches 5; Conservative 6; Mismatches 1; Indels 0; Gaps 0;

Db 80 gmcakcvcvdr1 91
   |:::| | | | | | | |
Qy 6 GTKSSKCVQKRV 17

RESULT 9
ENTRY S28969 #type complete
TITLE N-carbamoylserine amidohydrolase (EC 3.5.1.59) -
          Arthrobacter sp.
ORGANISM #formal name Arthrobacter sp.
DATE 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change
          19-Mar-1997

```

```
ACCESSIONS 528969
REFERENCE
#authors Romao, M.J.; Turk, D.; Comis-Rueth, F.X.; Huber, R.;
Schumacher, G.; Moellerling, H.; Ruessmann, L.
#journal J. Mol. Biol. (1992) 226:1111-1130
#title Crystal structure analysis, refinement and enzymatic reaction
mechanism of N-carbamoylisarcosine amidohydrolase from
Arthrobacter sp. at 2.0 A resolution.
#accession 528969
#status preliminary
#residues 1-264 ##label ROM
SUMMARY #length 264 #molecular-weight 29057 #checksum 6729

Query Match 40.4%; Score 57; DB 18; Length 264;
Best Local Similarity 40.0%; Pred. No. 2.22e+00;
Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Db 171 gataagcyrhtveda 185
| : : : ||| : || :
Qy 6 GTKSKCVRQKVEGS 20

RESULT 10
ENTRY B33485 #type complete
TITLE spore coat protein SP70 - slime mold (Dictyostelium
discoideum)
ORGANISM #formal name Dictyostelium discoideum
DATE 09-Mar-1990 #sequence_revision 11-Sep-1992 #text_change
30-Sep-1993
ACCESSIONS B33485
REFERENCE A33485
#authors Foaugh, K.L.; Loomis, W.F.
#journal Mol. Cell. Biol. (1989) 9:5215-5218
#title Spore coat genes SP60 and SP70 of Dictyostelium discoideum.
#cross-references MUID:90097939
#accession B33485
#status preliminary
#molecule_type DNA; mRNA
#residues 1-537 ##label FOS
#cross-references GB:M26238
#note the authors translated the codon AAT for residue 281 as
Aap
CLASSIFICATION #superfamily LDL receptor ligand-binding repeat homology
SUMMARY #length 537 #molecular-weight 56650 #checksum 2250

Query Match 39.7%; Score 56; DB 13; Length 537;
Best Local Similarity 54.5%; Pred. No. 3.42e+00;
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Db 291 kncgirdkve 301
| : : | : |||
Qy 8 KSKCVRQKVE 18

RESULT 11
ENTRY S21825 #type complete
TITLE vicilin-like storage protein G1b1-S, embryo - maize
ORGANISM #formal name Zea mays #common name maize
DATE 20-Feb-1995 #sequence_revision 20-Feb-1995 #text_change
20-Feb-1995
ACCESSIONS S21825
REFERENCE S21823
#authors Kriz, A.L.
#submission submitted to the EMBL Data Library, April 1991
```

```
#accession S21825
#status preliminary
#molecule_type DNA
#residues 1-540 ##label KRI
#cross-references EMBL:X59084
GENETICS
#gene G1b1-S
#introns 170/1; 195/2; 222/2; 319/2
SUMMARY #length 540 #molecular-weight 60239 #checksum 1419

Query Match 39.7%; Score 56; DB 13; Length 540;
Best Local Similarity 58.3%; Pred. No. 3.42e+00;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Db 32 hqghkagrcvrr 43
| | | : ||| :
Qy 4 HQGTKSKCVRQ 15

RESULT 12
ENTRY A53234 #type complete
TITLE globulin-1S, G1B1S - maize
ORGANISM #formal name Zea mays #common name maize
DATE 02-May-1994 #sequence_revision 18-Nov-1994 #text_change
05-Apr-1995
ACCESSIONS A53234
REFERENCE A53234
#authors Belanger, F.C.; Kriz, A.L.
#journal Genetics (1991) 129:863-872
#title Molecular basis for allelic polymorphism of the maize
Globulin-1 gene.
#cross-references MUID:92090707
#accession A53234
#status preliminary
#molecule_type DNA
#residues 1-573 ##label BEL
#cross-references NCBI:71280; NCBI:P:71284
#experimental_source inbred line Va 26
#note sequence extracted from NCBI backbone
SUMMARY #length 573 #molecular-weight 65075 #checksum 3569

Query Match 39.7%; Score 56; DB 13; Length 573;
Best Local Similarity 58.3%; Pred. No. 3.42e+00;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Db 32 hqghkagrcvrr 43
| | | : ||| :
Qy 4 HQGTKSKCVRQ 15

RESULT 13
ENTRY A32494 #type complete
TITLE transposable element Txc protein 1 - African clawed frog
ORGANISM #formal name Xenopus laevis #common name African clawed frog
DATE 12-Oct-1989 #sequence_revision 31-Dec-1993 #text_change
31-Dec-1993
ACCESSIONS A32494
REFERENCE A32494
#authors Garrett, J.E.; Knutzon, D.S.; Carroll, D.
#journal Mol. Cell. Biol. (1989) 9:3018-3027
#title Composite transposable elements in the Xenopus laevis genome.
#cross-references MUID:89384562
#accession A32494
#status preliminary
```

Jan 28 12:17

US-08-487-283A-1.rpr

11

```
#molecule type DNA
##residues 1-775 ##label GAR
##cross-references GB:M26915
##note the authors translated the codon ATT for residue as Gln,
and AAG for residue 288 as Leu
SUMMARY #length 775 #molecular-weight 82355 #checksum 6734

Query Match 39.0%; Score 55; DB 15; Length 775;
Best Local Similarity 46.7%; Pred. No. 5.22e+00;
Matches 7; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

Db 617 ntsskcvrkvvegss 631
Qy 7 TKSSKCVRKVRQKVE 21
::: ||| |||::
```

```
RESULT 14
ENTRY EA7119 #type complete
TITLE spore coat peptide CotZ - Bacillus subtilis
ORGANISM #formal_name Bacillus subtilis
DATE 21-Sep-1993 #sequence_revision 18-Nov-1994 #text_change
18-Nov-1994
ACCESSIONS EA7119
REFERENCE A47119
authors Zhang, J.; Fitz-James, P.C.; Aronson, A.I.
#journal J. Bacteriol. (1993) 175:3757-3766
#title Cloning and characterization of a cluster of genes encoding
polypeptides present in the insoluble fraction of the spore
coat of Bacillus subtilis.
#cross-references MUID:93285989
#accession EA7119
##status preliminary
##molecule type nucleic acid
##residues 1-148 ##label ZHA
#cross-references NCBIN:133538; NCIP:133548
#note sequence extracted from NCBI backbone
SUMMARY #length 148 #molecular-weight 16534 #checksum 4681
```

```
Query Match 38.3%; Score 54; DB 12; Length 148;
Best Local Similarity 63.6%; Pred. No. 7.93e+00;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Db 4 ktsscvreave 14
I:I |||: ||
Qy 8 KSSKCVRKQKVE 18
```

```
RESULT 15
ENTRY G64383 #type complete
TITLE riboflavin-specific deaminase (EC 3.5.4.-) - Methanococcus
jannaschii
ORGANISM #formal_name Methanococcus jannaschii
DATE 13-Sep-1996 #sequence_revision 13-Sep-1996 #text_change
13-Sep-1996
ACCESSIONS G64383
REFERENCE A64300
authors Bult, C.J.; White, O.; Olsen, G.J.; Zhou, L.; Fleischmann,
R.D.; Sutton, G.G.; Blake, J.A.; FitzGerald, L.M.; Clayton,
R.A.; Gocayne, J.D.; Kerlavage, A.R.; Dougherty, B.A.;
Tomb, J.F.; Adams, M.D.; Reich, C.I.; Overbeek, R.;
Kirkness, E.F.; Weinstock, K.G.; Merrick, J.M.; Glodek, A.;
Scott, J.L.; Geoghagan, N.S.M.; Weidman, J.F.; Fuhrmann,
J.L.; Nguyen, D.; Utterback, T.R.; Kelley, J.M.; Peterson,
J.D.; Sadow, P.W.; Hama, M.C.; Cotton, M.D.; Roberts,
```

Jan 28 12:17

US-08-487-283A-1.rpr

12

```
K.M.; Hurst, M.A.; Kaine, B.P.; Borodovsky, M.; Klenk,
H.P.; Fraser, C.M.; Smith, H.O.; Woese, C.R.; Venter, J.C.
#journal Science (1996) 273:1058-1073
#title Complete genome sequence of the methanogenic archaeon,
Methanococcus jannaschii.
#accession G64383
##status preliminary; nucleic acid sequence not shown;
translation not shown
##molecule type DNA
##residues 1-224 ##label BUL
#cross-references GB:L77117; TIGR:MJ0671; CDS_PID:g1510756
GENETICS
#map_position REV597638-596964
#start_codon TTG
KEYWORDS hydrolase
SUMMARY #length 224 #molecular-weight 25037 #checksum 2215
```

```
Query Match 38.3%; Score 54; DB 12; Length 224;
Best Local Similarity 33.3%; Pred. No. 7.93e+00;
Matches 6; Conservative 4; Mismatches 8; Indels 0; Gaps 0;

Db 118 iledmgvervkcgrgkv 135
::: | |||::
Qy 1 VIDHQTGTSKSKVRQKVE 18
```

Search completed: Wed Jan 28 12:10:33 1998  
Job time : 16 secs.



[4] RP SEQUENCE OF 678-751.  
RX MEDLINE: 79005687.  
RA FERNANDEZ H.N., HUGLI T.E.;  
RL J. BIOL. CHEM. 253:6955-6964(1978).  
[5] RP SEQUENCE OF 678-751 FROM N.A.  
RX MEDLINE: 91144547.  
RA BOHSACK J.F., MOLLISON K.W., BUKO A.M., ASHWORTH J.C., HILL H.R.;  
RL BIOCHEM. J. 273:635-640(1991).  
[6] RP STRUCTURE BY NMR OF C5A.  
RX MEDLINE: 88309754.  
RA ZUIDERWEG E.R., MOLLISON K.W., HENKIN J., CARTER G.W.;  
RL BIOCHEMISTRY 27:3568-3580(1988).  
[7] RP STRUCTURE BY NMR OF C5A.  
RX MEDLINE: 89207527.  
RA ZUIDERWEG E.R., NETTESHEIM D.G., MOLLISON K.W., CARTER G.W.;  
RL BIOCHEMISTRY 28:172-185(1989).  
[8] RP STRUCTURE BY NMR OF C5A.  
RX MEDLINE: 89274164.  
RA ZUIDERWEG E.R., FESIK S.W.;  
RL BIOCHEMISTRY 28:2387-2391(1989).  
CC -!- FUNCTION: ACTIVATION OF C5 BY A C5 CONVERTASE INITIATES THE  
CC SPONTANEOUS ASSEMBLY OF THE LATE COMPLEMENT COMPONENTS, C5-C9,  
CC INTO THE MEMBRANE ATTACK COMPLEX. C5B HAS A TRANSIENT BINDING SITE  
CC FOR C6. THE C5B-C6 COMPLEX IS THE FOUNDATION UPON WHICH THE LYTC  
CC COMPLEX IS ASSEMBLED.  
CC -!- SUBUNIT: C5 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 BASIC  
CC RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
CC BOND. C5 CONVERTASE ACTIVATES C5 BY CLEAVING THE ALPHA CHAIN,  
CC RELEASING C5A ANAPHYLATOXIN & GENERATING C5B (BETA CHAIN + ALPHA  
CC CHAIN).  
CC -!- SIMILARITY: TO C3, C4 AND ALPHA-2-MACROGLOBULIN.  
CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C5,  
CC C5 ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
CC INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
CC PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
CC BASOPHILIC LEUKOCYTES. C5A ALSO STIMULATES THE LOCOMOTION OF  
CC POLYMORPHONUCLEAR LEUKOCYTES (CHEMOKINESIS) AND DIRECT THEIR  
CC MIGRATION TOWARD SITES OF INFLAMMATION (CHEMOTAXIS).  
CC -!- CAUTION: REF. 3 SEQUENCE DIFFERS FROM THAT SHOWN FROM POSITION 855  
CC ONWARD DUE TO THE PRESENCE OF AN ALU REPEAT.  
CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.  
DR EMBL: M57729; G179983; --  
DR EMBL: M65134; G179692; --  
DR PIR: A40075; C5HU.  
DR PIR: S15121; S15121.  
DR HSP: P01032; IC5A.  
DR MIM: I20900; --  
DR PROSITE: PS00477; ALPHA\_2\_MACROGLOBULIN.  
KW COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; GLYCOPROTEIN;  
KW PLASMA; MEMBRANE ATTACK COMPLEX; CYTOLYSIS; INFLAMMATORY RESPONSE;  
KW SIGNAL; POLYMORPHISM.  
FT SIGNAL 1 18 POTENTIAL.  
FT CHAIN 19 673 COMPLEMENT C5 BETA CHAIN.  
FT PROPEP 674 677  
FT CHAIN 678 1676 COMPLEMENT C5 ALPHA CHAIN.  
FT PEPTIDE 678 751 C5A ANAPHYLATOXIN.  
FT CHAIN 752 1676 C5B (ALPHA').  
FT DOMAIN 698 732 ANAPHYLATOXIN-LIKE.  
FT DISULFID 698 724

FT DISULFID 699 731  
FT DISULFID 711 732  
FT CARBOHYD 741 741  
FT CARBOHYD 911 911 POTENTIAL.  
FT CARBOHYD 1115 1115 POTENTIAL.  
FT CARBOHYD 1630 1630 POTENTIAL.  
FT VARIANT 518 518 F -> S.  
SQ SEQUENCE 1676 AA; 188331 MW; 9D5C6E59 CRC32;  
  
Query Match 100.0%; Score 141; DB 2; Length 1676;  
Best Local Similarity 100.0%; Pred. No. 1.16e-24;  
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
Db 872 vidhgtksskcvrkvegs 892  
|||||  
Qy 1 VIDHGTSSKCVRKVEGSS 21  
  
RESULT 2  
ID C05 MOUSE STANDARD; PRT; 1680 AA.  
AC P06684;  
DT 01-JAN-1988 (REL. 06, CREATED)  
DT 01-NOV-1990 (REL. 16, LAST SEQUENCE UPDATE)  
DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)  
DE COMPLEMENT C5 PRECURSOR (CONTAINS: C5A ANAPHYLATOXIN).  
GN C5.  
OS MUS MUSCULUS (MOUSE).  
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;  
OC EUTHERIA; RODENTIA.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE: 90153853.  
RA WETSEL R.A., FLEISCHER D.T., HAVILAND D.L.;  
RL J. BIOL. CHEM. 265:2435-2440(1990).  
RN [2]  
RP SEQUENCE OF 41-1680 FROM N.A.  
RX MEDLINE: 87185363.  
RA WETSEL R.A., OGATA R.T., TACK B.F.;  
RL BIOCHEMISTRY 26:737-743(1987).  
CC -!- FUNCTION: ACTIVATION OF C5 BY A C5 CONVERTASE INITIATES THE  
CC SPONTANEOUS ASSEMBLY OF THE LATE COMPLEMENT COMPONENTS, C5-C9,  
CC INTO THE MEMBRANE ATTACK COMPLEX. C5B HAS A TRANSIENT BINDING SITE  
CC FOR C6. THE C5B-C6 COMPLEX IS THE FOUNDATION UPON WHICH THE LYTC  
CC COMPLEX IS ASSEMBLED.  
CC -!- SUBUNIT: C5 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 BASIC  
CC RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
CC BOND. C5 CONVERTASE ACTIVATES C5 BY CLEAVING THE ALPHA CHAIN,  
CC RELEASING C5A ANAPHYLATOXIN & GENERATING C5B (BETA CHAIN + ALPHA  
CC CHAIN).  
CC -!- SIMILARITY: TO C3, C4 AND ALPHA-2-MACROGLOBULIN.  
CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C5,  
CC C5 ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
CC INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
CC PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
CC BASOPHILIC LEUKOCYTES. C5A ALSO STIMULATES THE LOCOMOTION OF  
CC POLYMORPHONUCLEAR LEUKOCYTES (CHEMOKINESIS) AND DIRECT THEIR  
CC MIGRATION TOWARD SITES OF INFLAMMATION (CHEMOTAXIS).  
CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.  
DR EMBL: M35525; G309124; --  
DR EMBL: M35526; G309123; --  
DR PIR: A27538; A27538.  
DR PIR: A35530; A35530.  
DR HSP: P01032; IC5A.  
DR PROSITE: PS00477; ALPHA\_2\_MACROGLOBULIN.

KW COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; GLYCOPROTEIN;  
 KW PLASMA; MEMBRANE ATTACK COMPLEX; CYTOLYSIS; INFLAMMATORY RESPONSE;  
 KW SIGNAL.  
 FT SIGNAL 1 18  
 FT CHAIN 19 1680  
 FT CHAIN 19 674  
 FT PROPEP 675 678  
 FT CHAIN 679 1680  
 FT PEPTIDE 679 755  
 FT CHAIN 756 1680  
 FT DOMAIN 702 736  
 FT DISULFID 702 728  
 FT DISULFID 703 735  
 FT DISULFID 715 736  
 FT CARBOHYD 427 427  
 FT CARBOHYD 915 915  
 FT CARBOHYD 1119 1119  
 FT CARBOHYD 1633 1633  
 FT VARIANT 216 216  
 FT VARIANT 217 1680  
 SQ SEQUENCE 1680 AA; 188877 MW; AA17044B CRC32;

Query Match 48.9%; Score 69; DB 2; Length 1680;  
 Best Local Similarity 47.1%; Pred. No. 7.22e-04;  
 Matches 8; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

Db 880 hterparcvfgriegas 896  
 : ::::|::|::|::|::|  
 Qy 5 QTKSKCVRQKVEGSS 21

RESULT 3  
 ID IROA NEIME STANDARD; PRT; 943 AA.  
 AC Q06379;  
 DT 01-NOV-1995 (REL. 32, CREATED)  
 DT 01-NOV-1995 (REL. 32, LAST SEQUENCE UPDATE)  
 DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)  
 DE IRON-REGULATED OUTER MEMBRANE PROTEIN A PRECURSOR.  
 GN IROA.  
 OS NEISSERIA MENINGITIDIS.  
 OC PROKARYOTA; GRACILICUTES; SCOTOBACTERIA; AEROBIC RODS AND COCCI;  
 OC NEISSERIACEAE.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=BNVC;  
 RX MEDLINE; 94011384.  
 RA PETERSSON A., VAN DER LEY P., POOLMAN J.T., TOMMASSEN J.;  
 RL INFECT. IMMUN. 61:4724-4733(1993).  
 CC -!- FUNCTION: UNKNOWN. MAY BE AN IRON-SIDEROPHORE RECEPTOR.  
 CC -!- SUBCELLULAR LOCATION: OUTER MEMBRANE.  
 CC -!- INDUCTION: BY IRON-STARVATION CONDITIONS.  
 CC -!- SIMILARITY: LOCAL TO OTHER TONB-DEPENDENT RECEPTOR PROTEINS.  
 DR EMBL; X69214; G45064; -.  
 DR PROSITE; PS00430; TONB DEPENDENT REC 1.  
 DR PROSITE; PS01156; TONB DEPENDENT REC 2.  
 KW OUTER MEMBRANE; IRON TRANSPORT; TONB BOX; SIGNAL; RECEPTOR.  
 FT SIGNAL 1 27  
 FT CHAIN 28 943  
 FT CHAIN 826 943  
 SQ SEQUENCE 943 AA; 105424 MW; 16644948 CRC32;

Query Match 41.8%; Score 59; DB 5; Length 943;  
 Best Local Similarity 50.0%; Pred. No. 1.65e-01;  
 Matches 7; Conservative 5; Mismatches 2; Indels 0; Gaps 0;

Db 595 tsrkcvprkingsn 608  
 :|::|::|::|::|::|::|::|  
 Qy 8 KSKCVRQKVEGSS 21

RESULT 4  
 ID YEMA DROME STANDARD; PRT; 1002 AA.  
 AC P25992;  
 DT 01-MAY-1992 (REL. 22, CREATED)  
 DT 01-MAY-1992 (REL. 22, LAST SEQUENCE UPDATE)  
 DT 01-FEB-1995 (REL. 31, LAST ANNOTATION UPDATE)  
 DE YEMANUCLEIN-ALPHA.  
 GN YEMA OR YG4.5.  
 OS DROSOPHILA MELANOGASTER (FRUIT FLY).  
 OC EUKARYOTA; METAZOA; ARTHROPODA; INSECTA; DIPTERA.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=CANTON-S;  
 RX MEDLINE; 92297435.  
 RA AIT-AHMED O., BELLON B., CAPRI M., JOBLET C., THOMAS-DELAAGE M.;  
 RL MECH. DEV. 37:69-80(1992).  
 CC -!- FUNCTION: MAY PLAY A KEY ROLE IN EGG ORGANIZATION. IT MAY BE A  
 CC TRANSCRIPTIONAL REGULATOR.  
 CC -!- PTM: THE N-TERMINAL IS BLOCKED.  
 CC -!- TISSUE SPECIFICITY: OOCYTE-SPECIFIC.  
 CC -!- DEVELOPMENTAL STAGE: EXPRESSED AT ALL OOGENIC STAGES.  
 CC -!- SUBCELLULAR LOCATION: NUCLEAR.  
 DR EMBL; X63503; G8838; -.  
 DR PIR; S22146; S22146.  
 DR FLYBASE; FBGN005596; YEM-ALPHA.  
 KW NUCLEAR PROTEIN; DNA-BINDING; REPEAT.  
 FT DOMAIN 80 85  
 FT DOMAIN 207 217  
 FT DOMAIN 219 261  
 FT DOMAIN 230 253  
 FT REPEAT 230 241  
 FT REPEAT 242 253  
 FT VARIANT 698 698  
 SQ SEQUENCE 1002 AA; 109310 MW; 955FD2C1 CRC32;

Query Match 41.8%; Score 59; DB 11; Length 1002;  
 Best Local Similarity 50.0%; Pred. No. 1.65e-01;  
 Matches 6; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

Db 47 tktakcirikid 58  
 |::|::|::|::|::|::|::|  
 Qy 7 TKSKCVRQKVE 18

RESULT 5  
 ID RL34 HUMAN STANDARD; PRT; 116 AA.  
 AC P49207;  
 DT 01-FEB-1996 (REL. 33, CREATED)  
 DT 01-FEB-1996 (REL. 33, LAST SEQUENCE UPDATE)  
 DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)  
 DE 60S RIBOSOMAL PROTEIN L34.  
 GN RPL34.  
 OS HOMO SAPIENS (HUMAN).  
 OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;  
 OC EUTHERIA; PRIMATES.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-OVARY;

Jan 28 12:17

US-08-487-283A-1.fsp

7

RX MEDLINE; 96039267.  
RA ROMMENS J.M., DUROCHER F., MCARTHUR J., TONIN P., LEBLANC J.F.,  
RA ALLEN T., SAMSON C., FERRI L., NAROD S., MORGAN K., SIMARD J.;  
RL GENOMICS 28:530-542(1995).  
CC -!- SIMILARITY: BELONGS TO THE L34E FAMILY OF RIBOSOMAL PROTEINS.  
DR EMBL; L38941; G1008856; -.  
KW RIBOSOMAL PROTEIN.  
FT INIT MET 0 0 BY SIMILARITY.  
SQ SEQUENCE 116 AA; 13174 MW; 490F4AF1 CRC32;

Query Match 40.4%; Score 57; DB 8; Length 116;  
Best Local Similarity 41.7%; Pred. No. 4.57e-01;  
Matches 5; Conservative 6; Mismatches 1; Indels 0; Gaps 0;

Db 79 gmcakcvrdri 90  
|:::|||||::: 17  
Qy 6 GTKSSKVRQKV 17

RESULT 6  
ID GSH ARTSP STANDARD; PRT; 264 AA.  
AC P32400;  
DT 01-OCT-1993 (REL. 27, CREATED)  
DT 01-OCT-1993 (REL. 27, LAST SEQUENCE UPDATE)  
DT 01-FEB-1995 (REL. 31, LAST ANNOTATION UPDATE)  
DE N-CARBAMOYL-SARCOSE AMIDASE (EC 3.5.1.59) (N-CARBAMOYL-SARCOSE  
DE AMIDOHYDROLASE) (CSHASE).  
OS ARTHROBACTER SP.  
OC PROKARYOTA; FIRMICUTES; IRREGULAR ASPOGENOUS RODS; CORYNEFORM GROUP.  
RN [1]  
RP X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS), AND REVISIONS TO 184 AND 232.  
RX MEDLINE; 92389321.  
RA ROMAO M.J., TURK D., GOMIS-RUETH F.-X., HUBER R.;  
RL J. MOL. BIOL. 226:1111-1130(1992).  
CC -!- CATALYTIC ACTIVITY: N-CARBAMOYL-SARCOSE + H(2)O = SARCOSE +  
CO(2) + NH(3).  
CC -!- SUBUNIT: HOMOTETRAMER.  
CC -!- COFACTOR: ONE SULFATE ION PER SUBUNIT.  
CC -!- PATHWAY: DEGRADATION OF CREATININE TO GLYCINE.  
DR PIR; S28969; S28969.  
DR PDB; INBA; 22-JUN-94.  
KW HYDROLASE; 3D-STRUCTURE.  
FT ACT SITE 177 177 INVOLVED IN HYDROLYSIS OF THE SUBSTRATE.  
SQ SEQUENCE 264 AA; 29057 MW; 81A56865 CRC32;

Query Match 40.4%; Score 57; DB 2; Length 264;  
Best Local Similarity 40.0%; Pred. No. 4.57e-01;  
Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Db 171 gataagcvrhtvda 185  
|:::|||||::: 20  
Qy 6 GTKSSKVRQKVEGS 20

RESULT 7  
ID SP70 DICDI STANDARD; PRT; 537 AA.  
AC P15269; P08126;  
DT 01-AUG-1988 (REL. 08, CREATED)  
DT 01-APR-1990 (REL. 14, LAST SEQUENCE UPDATE)  
DT 01-FEB-1994 (REL. 28, LAST ANNOTATION UPDATE)  
DE SPORE COAT PROTEIN SP70 PRECURSOR (BEEJIN PROTEIN).  
GN COYB.  
OS DICTYOSTELIUM DISCOIDEUM (SLIME MOLD).  
CC EUKARYOTA; PROTOZOA; SARCOMASTIGOPHORA; SARCODINA; RHIZOPODA;

Jan 28 12:17

US-08-487-283A-1.fsp

8

OC EUMYCETOZOA; DICTYOSTELIA.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE; 90097939.  
RA FOSNAUGH K.L., LOOMIS W.F.;  
RL MOL. CELL. BIOL. 9:5215-5218(1989).  
RN [2]  
RP PRELIMINARY SEQUENCE OF 72-170 FROM N.A.  
RX MEDLINE; 87057653.  
RA GOMER R.H., DATTA S., FIRTEL R.A.;  
RL J. CELL BIOL. 103:1999-2015(1986).  
DR EMBL; M26238; G167889; -.  
DR PIR; B33485; B33485.  
DR PIR; B25439; B25439.  
DR DICTYDB; DD03009; COYB.  
KW GLYCOPROTEIN; PHOSPHORYLATION; REPEAT; SPORULATION; SIGNAL.  
FT SIGNAL 1 20  
FT CHAIN 21 537 SPORE COAT PROTEIN SP70.  
FT DOMAIN 182 250 SER/THR-RICH.  
FT DOMAIN 190 248 5.5 X 11 AA TANDEM REPEATS.  
FT REPEAT 190 200 1.  
FT REPEAT 191 211 2.  
FT REPEAT 212 222 3.  
FT REPEAT 223 233 4.  
FT REPEAT 234 244 5.  
FT REPEAT 245 248 6 (INCOMPLETE).  
FT REPEAT 256 263 PRESPORE MOTIF.  
FT REPEAT 284 291 PRESPORE MOTIF.  
FT REPEAT 364 371 PRESPORE MOTIF.  
FT CARBOHYD 97 POTENTIAL.  
SQ SEQUENCE 537 AA; 56650 MW; 5D59CBAC CRC32;

Query Match 39.7%; Score 56; DB 9; Length 537;  
Best Local Similarity 54.5%; Pred. No. 7.54e-01;  
Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Db 291 kngecirldkve 301  
|:::|:|:|:|  
Qy 8 KSKKVRQKVE 18

RESULT 8  
ID GLB1 MAIZE STANDARD; PRT; 573 AA.  
AC P15590;  
DT 01-APR-1990 (REL. 14, CREATED)  
DT 01-AUG-1990 (REL. 15, LAST SEQUENCE UPDATE)  
DT 01-OCT-1994 (REL. 30, LAST ANNOTATION UPDATE)  
DE GLOBULIN-1 S ALLELE PRECURSOR (GLB1-S) (7S-LIKE).  
GN GLB1.  
OS ZEA MAYS (MAIZE).  
OC EUKARYOTA; PLANTA; EMBRYOPHYTA; ANGIOSPERMAE; MONOCOTYLEDONEAE;  
OC CYPERALES; GRAMINEAE.  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN-CV. INBRED LINE VA26;  
RA BELANGER F.C., KRIZ A.L.;  
RL PLANT PHYSIOL. 91:636-643(1989).  
RN [2]  
RP SEQUENCE OF 87-100.  
RX MEDLINE; 89374022.  
RA KRIZ A.L.;  
RL BIOCHEM. GENET. 27:239-251(1989).  
CC -!- SIMILARITY: TO OTHER 7S SEED STORAGE PROTEINS (PHASEOLIN, VICILIN,  
CONVICILIN, CONGLYCININ, ETC.).

CC -!- POLYMORPHISM: THE THREE MOST COMMONLY OCCURRING GIBI ALLELES HAVE  
CC THE DESIGNATION L, I, AND S FOR LARGE, INTERMEDIATE, AND SMALL  
CC PROTEINS, RESPECTIVELY.

CC -!- PTM: THREE PROTEIN-PROCESSING STEPS OCCUR IN THE FORMATION OF THE  
CC MATURE PROTEIN FROM THE PRIMARY TRANSLATION PRODUCT.

DR EMBL; M24845; G168481; -.

DR HSP; P02853; 1CAU.

DR MAIZEDB; 30181; -.

KW SEED STORAGE PROTEIN; SIGNAL.

FT SIGNAL 1 18 OR 21 (POTENTIAL).

FT PROPEP 19 86

FT CHAIN 87 573 GLOBULIN-1 S.

FT CARBOHYD 349 349 POTENTIAL.

SQ SEQUENCE 573 AA; 65029 MW; 7E755E20 CRC32;

Query Match 39.7%; Score 56; DB 4; Length 573;

Best Local Similarity 58.3%; Pred. No. 7.54e-01;

Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Db 32 hghkgcgvrr 43

|||||

Qy 4 HQTGKSKVRQ 15

RESULT 9

ID RPOB PSEPU STANDARD; PRT; 1357 AA.

AC P19175;

DT 01-NOV-1990 (REL. 16, CREATED)

DT 01-MAR-1992 (REL. 21, LAST SEQUENCE UPDATE)

DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)

DE DNA-DIRECTED RNA POLYMERASE BETA CHAIN (EC 2.7.7.6) (TRANSCRIPTASE

DE BETA CHAIN) (RNA POLYMERASE BETA SUBUNIT).

GN RPOB.

OS PSEUDOMONAS PUTIDA.

OC PROKARYOTA; GRACILICUTES; SCOTOBACTERIA; AEROBIC RODS AND COCCI;

OC PSEUDOMONADACEAE.

RN [1]

RP SEQUENCE FROM N.A.

RA BORODIN A.M., DANILKOVICH A.V., ALLIKMETS R.L., ROSTAPSHOV V.M.,

RA CHERNOV I.P., AZHIKINA T.L., MONASTYRSKAYA S., SVERDLOV D.;

RL DOKL. BIOCHEM. 302:1261-1265(1988).

RN [2]

RP SEQUENCE OF 1036-1357 FROM N.A.

RX MEDLINE; 89117617.

RA BORODIN A.M., DANILKOVICH A.V., CHERNOV I.I., AZHIKINA T.L.,

RA ROSTAPSHOV V.M., MONASTYRSKAYA G.S.;

RL BIOORG. KHIM. 14:1179-1182(1988).

CC -!- FUNCTION: DNA-DEPENDENT RNA POLYMERASE CATALYZES THE TRANSCRIPTION

CC OF DNA INTO RNA USING THE FOUR RIBONUCLEOSIDE TRIPHOSPHATES AS

CC SUBSTRATES.

CC -!- CATALYTIC ACTIVITY: N NUCLEOSIDE TRIPHOSPHATE = N PYROPHOSPHATE +

CC RNA (N).

CC -!- SUBUNIT: THE ENZYME CONSISTS OF THE SIGMA CHAIN AND THE CORE

CC ENZYME WHICH IS COMPOSED OF 2 ALPHA CHAINS, 1 BETA CHAIN, AND 1

CC BETA' CHAIN.

CC -!- SIMILARITY: BELONGS TO THE RNA POLYMERASE BETA CHAIN FAMILY.

DR EMBL; X15849; G45729; -.

DR EMBL; M38319; G151547; -.

KW TRANSCRIPTION; DNA-DIRECTED RNA POLYMERASE.

FT CONFLICT 1180 1180 T -> N (IN REF. 2).

FT CONFLICT 1184 1184 I -> V (IN REF. 2).

FT CONFLICT 1236 1236 F -> S (IN REF. 2).

SQ SEQUENCE 1357 AA; 151305 MW; BBF88A37 CRC32;

Query Match 39.7%; Score 56; DB 8; Length 1357;

Best Local Similarity 41.2%; Pred. No. 7.54e-01;

Matches 7; Conservative 4; Mismatches 6; Indels 0; Gaps 0;

Db 450 idhlgrrvrcvcmgmae 466

|||||

Qy 2 IDHGTGKSKVRQKVE 18

RESULT 10

ID YTX1 XENLA STANDARD; PRT; 775 AA.

AC P14380;

DT 01-JAN-1990 (REL. 13, CREATED)

DT 01-JAN-1990 (REL. 13, LAST SEQUENCE UPDATE)

DT 01-APR-1990 (REL. 14, LAST ANNOTATION UPDATE)

DE TRANSPOSON TX1 HYPOTHETICAL 82 KD PROTEIN (ORF 1).

OS XENOPUS LAEVIS (AFRICAN CLAWED FROG).

OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; AMPHIBIA; ANURA.

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE; 89384562.

RA GARRETT J.E., KNUZZON D.S., CARROLL D.;

RL MOL. CELL. BIOL. 9:3018-3027(1989).

DR EMBL; M26915; G214845; -.

DR PIR; A32494; A32494.

KW HYPOTHETICAL PROTEIN; TRANSPOSABLE ELEMENT.

SQ SEQUENCE 775 AA; 82355 MW; 9738B05A CRC32;

Query Match 39.0%; Score 55; DB 11; Length 775;

Best Local Similarity 46.7%; Pred. No. 1.24e+00;

Matches 7; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

Db 617 sntekcvsvsevegt 631

|||||

Qy 7 TKSSKCVRKVEGSS 21

RESULT 11

ID COTZ B-ACSU STANDARD; PRT; 148 AA.

AC Q08312;

DT 01-OCT-1994 (REL. 30, CREATED)

DT 01-OCT-1994 (REL. 30, LAST SEQUENCE UPDATE)

DT 01-FEB-1995 (REL. 31, LAST ANNOTATION UPDATE)

DE SPORE COAT PROTEIN 2.

GN COTZ.

OS BACILLUS SUBTILIS.

OC PROKARYOTA; FIRMICUTES; ENDOSPORE-FORMING RODS AND COCCI; BACILLACEAE.

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=168 / JH642;

RX MEDLINE; 93285989.

RA ZHANG J., FITZ-JAMES P.C., ARONSON A.I.;

RL J. BACTERIOL. 175:3757-3766(1993).

CC -!- SUBCELLULAR LOCATION: SPORE OUTER COAT.

CC -!- SUBUNIT: DISULFIDE CROSS-LINKED EITHER TO ITSELF OR TO COTY.

CC -!- SIMILARITY: TO COTY.

DR EMBL; L10116; G304149; -.

DR PIR; E47119; E47119.

DR SUBTILIST; BG10500; COTZ.

KW SPORULATION.

SQ SEQUENCE 148 AA; 16534 MW; B5442F5E CRC32;

Query Match 38.3%; Score 54; DB 2; Length 148;

Best Local Similarity 63.6%; Pred. No. 2.01e+00;





Jan 28 12:17

US-08-487-283A-1.FSP

13

RX MEDLINE; 83164198.  
RA SARNOW P., HEARING P., ANDERSON C.W., REICH N., LEVINE A.J.;  
RL J. MOL. BIOL. 162:565-583(1982).  
RN [2]  
RP COMPLETE GENOME.  
RX MEDLINE; 92087470.  
RA CHROBOCZEK J., BIEBER F., JACROT B.;  
RL VIROLOGY 186:280-285(1992).  
DR EMBL; M73260; -: NOT\_ANNOTATED\_CDS.  
DR EMBL; X02998; G58502; -.  
DR PIR; B03807; Q4ADE5.  
KW EARLY PROTEIN.  
SQ SEQUENCE 116 AA; 13298 MW; 66EA9B5C CRC32;  
  
Query Match 37.6%; Score 53; DB 3; Length 116;  
Best Local Similarity 60.0%; Pred. No. 3.25e+00;  
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;  
  
Db 3 rclrlkvega 12  
:|:| | | | |  
Qy 11 KCVRKVEGS 20

RESULT 15  
ID E411 ADE02 STANDARD; PRT; 116 AA.  
AC P03241;  
DT 21-JUL-1986 (REL. 01, CREATED)  
DT 21-JUL-1986 (REL. 01, LAST SEQUENCE UPDATE)  
DT 01-APR-1990 (REL. 14, LAST ANNOTATION UPDATE)  
DE PROBABLE EARLY EA 11 KD PROTEIN.  
OS HUMAN ADENOVIRUS TYPE 2.  
OC VIRIDAE; DS-DNA NONENVELOPED VIRUSES; ADENOVIRIDAE; MASTADENOVIRUSES.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE; 82059444.  
RA HERRISE J., RIGOLET M., DUPONT DE DINECHIN S., GALIBERT F.;  
RL NUCLEIC ACIDS RES. 9:4023-4042(1981).  
DR EMBL; J01917; G209839; -.  
DR PIR; A03807; Q4ADE2.  
KW EARLY PROTEIN.  
SQ SEQUENCE 116 AA; 13255 MW; 950D6981 CRC32;

Query Match 37.6%; Score 53; DB 3; Length 116;  
Best Local Similarity 60.0%; Pred. No. 3.25e+00;  
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;  
  
Db 3 rclrlkvega 12  
:|:| | | | |  
Qy 11 KCVRKVEGS 20

Search completed: Wed Jan 28 12:11:02 1998  
Job time : 10 secs.

Release 2.1D John F. Collins, Biocomputing Research Unit.  
Copyright (c) 1993, 1994, 1995 University of Edinburgh, U.K.  
Distribution rights by IntelliGenetics, Inc.

Run on: Wed Jan 28 12:09:44 1998; MasPar time 2.31 Seconds  
111.852 Million cell updates

Tabular output not generated.

```
>US-08-487-283A-1
Title:
Description: (1-21) from US08487283A, pep
Perfect Score: 141
Sequence: 1 VIDHGTGSSKCVROKVESS 21
```

Scoring table: PAM 150

Searched: 101610 seqs. 12294212 residues

Post-processing: Minimum Match 0%  
Listing first 45 summaries

Database: a-geneseq28  
1:part1 2:part2 3:part3 4:part4 5:part5 6:part6 7:part7  
8:part8 9:part9 10:part10 11:part11 12:part12 13:part13  
14:part14 15:part15 16:part16 17:part17 18:part18  
19:part19 20:part20 21:part21

Statistics: Mean 18.854; Variance 56.108; scale 0.336

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Result No.	Score	Query Match	Length	DB	ID	Description	Pred. No.
1	141	100.0	21	15	R7605	Pro-C5 polypeptide KS	1.23e-09
2	141	100.0	1676	15	R7604	Pro-C5 polypeptide.	1.23e-09
3	55	39.0	264	4	R2271	CSHase.	3.50e+01
4	54	38.3	1732	17	R9629	P. gingivalis porphyr	4.47e+01
5	54	38.3	3163	16	R94347	Hepatitis GB virus (H	4.47e+01
6	53	37.6	589	3	R14327	Mouse epithelin precu	5.69e+01
7	53	37.6	652	15	R88124	Tobacco mosaic virus	5.69e+01
8	53	37.6	1009	5	R26206	Type B human platelet	5.69e+01
9	53	37.6	1089	2	R06910	Alpha type PDGF recep	5.69e+01
10	53	37.6	1089	2	R08267	Platelet derived grow	5.69e+01
11	53	37.6	1196	19	W04326	Rat petrin.	5.69e+01

## ALIGNMENTS

RESULT	1	
AC	R77605; standard; Protein; 21 AA.	
AC	R77605;	
DT	02-APR-1996 (first entry)	
DE	Pro-C5 polypeptide KSSKC epitope.	
KW	Complement C5; haemolysis; kidney; glomerulonephritis;	
KW	monoclonal antibody; antiinflammatory; antibody engineering;	
KW	humanised antibody; KSSKC epitope.	
OS	homo sapiens.	
PN	W09529697-A1.	
PN	W09529697-A1.	
PD	09-NOV-1995.	
PF	01-MAY-1995; U05688.	
PF	02-MAY-1994; U5-236208.	
PA	(ALEX-) ALEXION PHARM INC.	
PI	Evans MJ, Matis L, Mueller EE, Nye SH, Rollins S;	
PI	Rother RP, Springhorn J P, Squinto SP, Thomas TC;	
PI	Wang Y, Wilkins JA;	
PI	WPI; 95-392923/50.	
PR	Treating glomerulonephritis with antibody complement C5	
PT	component - to inhibit complement induced cell lysis	
PT	Example 13; Page 81; 181pp; English.	
PS	The cDNA sequence of the complement C5 gene transcript predicts a	
CCC	secreted pro-C5 precursor of 1676 amino acids (R77604). C5 is a	
CCC	beta-globulin heterodimer thought to play a role in the pathogenesis	
CCC	of glomerulonephritis (GN). Cleavage of the C5 alpha-chain	
CCC	by a convertase enzyme generates anaphylatoxic C5a. Monoclonal	

Jan 28 12:16

US-08-487-283A-1.rag

3

CC and humanised recombinant antibodies that recognise the alpha-chain  
CC KSKC epitope (R77605) block C5a generation, thereby reducing  
CC glomerular inflammation and kidney dysfunction associated with GN.  
SQ Sequence 21 AA;

Query Match 100.0%; Score 141; DB 15; Length 21;  
Best Local Similarity 100.0%; Pred. No. 1.23e-09;  
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 vidhgktkaskcvrkvqvegs 21  
|||||  
Qy 1 VIDHQGTSSKCKVRQKVEGSS 21

RESULT 2  
ID R77604 standard; Protein; 1676 AA.  
AC R77604;  
DT 15-MAR-1996 (first entry)  
DE Pro-C5 polypeptide.  
KW Complement C5; haemolysis; kidney; glomerulonephritis;  
KW monoclonal antibody; antiinflammatory; antibody engineering;  
KW humanised antibody.  
OS Homo sapiens.  
FH Key Location/Qualifiers  
FT Peptide 1..18  
FT /label= Sig\_peptide  
FT Protein 19..673  
FT /label= Beta-chain  
FT Cleavage\_site 673..674  
FT Cleavage\_site 674..678  
FT Peptide 674..677  
FT /label= Cleavage\_peptide  
FT Protein 678..1676  
FT /label= Alpha-chain  
FT /note= "amino acids 872-892 (854-874 of  
FT the mature protein) comprise the KSKS  
FT epitope".  
FT Peptide 678..751  
FT /label= C5a  
FT Cleavage\_site 751..752  
FT /label= Convertase cleavage\_site  
FT Modified\_site 911  
FT /label= N-glycosylation\_site  
FT Modified\_site 1115  
FT /label= N-glycosylation\_site  
FT Modified\_site 1630  
FT /label= N-glycosylation\_site  
FT W09529697-A1.  
PD 09-NOV-1995.  
PF 01-MAY-1995; U05688.  
PR 02-MAY-1994; US-236208.  
PA (ALEX-) ALEXION PHARM INC.  
PI Evans MJ, Matlis L, Mueller EE, Nye SH, Rollins S;  
PI Rother RP, Springhorn J P, Squinto SP, Thomas TC;  
PI Wang Y, Wilkins JA;  
DR WPI; 95-392923/50.  
PT Treating glomerulonephritis with antibody against complement C5  
PT component - to inhibit complement induced cell lysis  
PS Example 13; Page 82-92; 181pp; English.  
CC The cDNA sequence of the complement C5 gene transcript predicts a  
CC secreted pro-C5 precursor of 1676 amino acids (R77604). C5 is a  
CC beta-globulin heterodimer thought to play a role in the pathogenesis  
CC of glomerulonephritis (GN). Cleavage of the C5 alpha-chain  
CC by a convertase enzyme generates anaphylatoxic C5a. Monoclonal

Jan 28 12:16

US-08-487-283A-1.rag

4

CC and humanised recombinant antibodies that recognise the alpha-chain  
CC KSKC epitope (R77605) block C5a generation, thereby reducing  
CC glomerular inflammation and kidney dysfunction associated with GN.  
SQ Sequence 1676 AA;

Query Match 100.0%; Score 141; DB 15; Length 1676;  
Best Local Similarity 100.0%; Pred. No. 1.23e-09;  
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 872 vidhgktkaskcvrkvqvegs 892  
|||||  
Qy 1 VIDHQGTSSKCKVRQKVEGSS 21

RESULT 3  
ID R22271 standard; Protein; 264 AA.  
AC R22271;  
DT 30-JUL-1992 (first entry)  
DE CSHase.  
KW N-carbamoyl-sarcosine amidohydrolase; CSH; assay; diagnosis;  
KW creatinine.  
OS Arthrobacter sp. DSM 2563.  
PN EP-476670-A.  
PD 25-MAR-1992.  
PF 19-SEP-1991; 115974.  
PR 20-SEP-1990; DE-029844.  
PA (BOEF) BOEHRINGER MANNHEIM GMBH.  
PI Butscher H, Schumacher G;  
DR WPI; 92-098378/13.  
DR N-PSDB; Q22713.  
PT Recombinant DNA encoding N-carbamoyl-sarcosine-amidohydrolase -  
PT useful in clinical assay of creatinine, and vectors providing  
PT efficient expression in E.coli  
PS Claim 9; Page 9 + 7; 12pp; German.  
CC The sequence encoding CSHase is useful in assay of creatinine  
CC (for diagnosis of kidney disease). It can now be prepd. more  
CC simply than by known methods which involve culture of Arthrobacter  
CC on complex media.  
SQ Sequence 264 AA;

Query Match 39.0%; Score 55; DB 4; Length 264;  
Best Local Similarity 46.2%; Pred. No. 3.50e+01;  
Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Db 171 gataagcvrhtve 183  
|: :|: |  
Qy 6 GTKSSKCVRQKVE 18

RESULT 4  
ID R96029 standard; Protein; 1732 AA.  
AC R96029;  
DT 04-SEP-1996 (first entry)  
DE P. gingivalis porphyra.  
KW Porphyra; haemagglutinin; periodontal disease; vaccine; antibody.  
OS Porphyromonas gingivalis strain W12.  
FH Key Location/Qualifiers  
FT Region 688..708  
FT /note= "Pro-Asn repeat region type 1"  
FT Region 887..952  
FT /note= "Pro-Asn repeat region type 2"  
FT Region 946..967  
FT /note= "Pro-Asn repeat region type 1"  
FT Region 985..1006

PF 14-FEB-1995; U02118.  
PR 14-FEB-1994; US-196030.  
PR 13-MAY-1994; US-242654.  
PR 29-JUL-1994; US-283314.

Key	Location/Qualifiers
FFH	
FT	1..589
FT	Protein
FT	/label= precursor
FT	/note= "claim 21, page 55"
FT	280..335
FT	Protein
FT	/label= EP-1
FT	/note= "claim 22, page 55"
FT	205..261
FT	Protein
FT	/label= EP-2
FT	/note= "claim 23, page 55"
FT	59..114
FT	Peptide
FT	/label= EP
FT	/note= "claim 24, page 55"
FT	123..179
FT	Peptide
FT	/label= EP
FT	/note= "claim 25, page 55"
FT	362..416
FT	Peptide
FT	/label= EP
FT	/note= "claim 26, page 56"
FT	440..495
FT	Peptide
FT	/label= EP
FT	/note= "claim 27, page 56"
FT	515..570
FT	Peptide
FT	/label= EP

FT /note= \*claim 28, page 56\*  
 PN W09115510-A.  
 PD 17-OCT-1991.  
 PF 03-APR-1991; U02321.  
 PR 03-APR-1990; US-504508.  
 PR 13-MAR-1991; US-083796.  
 PA (BRIM ) BRISTOL-MYERS SQUIB.  
 PI Shoyab M, Plowman GD;  
 DR WPI; 91-325168/44.  
 DR N-PSDB; Q14340.  
 PT New cysteine-rich growth modulating proteins, epithelins - useful  
 PT as inhibitors of neoplastic cell growth and to promote wound  
 PT healing and treat psoriasis  
 PS Disclosure; Fig 23; 97pp; English.  
 CC ET-1 is a bifunctional growth regulator, capable of stimulating  
 CC the growth of some cell types while inhibiting the growth of others.  
 CC ET-2 is functionally similar to ET-1 w.r.t. growth inhibitory  
 CC bioactivity. In contrast, however, ET-2 is apparently not capable of  
 CC eliciting the growth stimulatory activity characteristic of ET-1 and,  
 CC in fact, antagonises this ET-1 activity.  
 CC See also Q14338-40, Q14952-53, R14328-9 and R15315-20.  
 SQ Sequence 589 AA;

Query Match 37.6%; Score 53; DB 3; Length 589;  
 Best Local Similarity 35.7%; Pred. No. 5.69e+01;  
 Matches 5; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

Db 560 hcsargtkclrkki 573

Qy | : : : : : | : | :

Qy 4 HQGTSKSKVRQKV 17

## RESULT 7

ID R88124 standard; Protein; 652 AA.  
 AC R88124;  
 DT 28-MAR-1996 (first entry)  
 DE Tobacco mosaic virus resistance N gene truncated protein.  
 KW Tobacco mosaic virus resistance; TMV; N gene; Solanaceae;  
 KW crop improvement; transgenic plant; crop improvement.  
 OS Nicotiana glauca.  
 PN W0935024-A1.  
 PD 28-DEC-1995.  
 PF 16-JUN-1995; U07754.  
 PR 17-JUN-1994; US-261663.  
 PA (RECC ) UNIV CALIFORNIA.  
 PA (USDA ) US SEC OF AGRIC.  
 PI Baker BJ, Whitham SA;  
 DR WPI; 96-058144/06.  
 DR N-PSDB; T09342.  
 PT Plant virus resistance gene N sequences from tobacco - useful for  
 PT generating transgenic Solanaceous plants resistant to Tobacco Mosaic  
 PT Virus  
 PS Claim 28; Page 75-79; 98pp; English.  
 CC The Nicotiana glauca N gene truncated protein (R88124) mediates  
 CC resistance to tobacco mosaic virus (TMV). A cDNA clone (T09342)  
 CC coding for the protein was obtd. from a N. glauca leaf cDNA  
 CC library by transposon tagging. DNA sequences encoding the  
 CC protein can be used to generate transgenic plants, esp. Solanaceae,  
 CC resistant to TMV.  
 SQ Sequence 652 AA;

Query Match 37.6%; Score 53; DB 15; Length 652;  
 Best Local Similarity 31.3%; Pred. No. 5.69e+01;  
 Matches 5; Conservative 5; Mismatches 6; Indels 0; Gaps 0;

Db 156 dnrdctdadcirqv 171  
 | : : : | : | : |  
 Qy 3 DHQGTSSKVRQKVE 18

## RESULT 8

ID R26206 standard; Protein; 1009 AA.  
 AC R26206;  
 DT 09-FEB-1993 (first entry)  
 DE Type B human platelet-derived growth factor receptor.  
 KW PDGF; PDGF-R; mesenchyme; tyrosine kinase; ligand binding region.  
 OS Homo sapiens.  
 FH Key Location/Qualifiers  
 FT Peptide 1..23  
 FT /label= Signal\_peptide  
 FT Protein 24..1009  
 FT /label= Mature\_PDGF-A  
 PN W09213867-A.  
 PD 20-AUG-1992.  
 PF 28-JAN-1992; U00730.  
 PR 31-JAN-1991; US-650793.  
 PA (CORTE-) COR THERAPEUTICS INC.  
 PI Escobedo JA, Fretto LJ, Giese NA, Tomlinson JE, Williams LT;  
 PI Wolf D;  
 DR WPI; 92-299970/36.  
 DR N-PSDB; Q27451.  
 PT Platelet derived growth factor receptor (PDGF-R) poly:peptide(s)  
 PT - useful as therapeutic and diagnostic agents e.g. for assaying  
 PT PDGF activity in sample  
 PS Disclosure; Page 90; 109pp; English.  
 CC The sequence given is one allele of type A human platelet-derived  
 CC growth factor (PDGF) receptor (PDGF-R). This receptor is typically  
 CC found on cells of mesenchymal origin. It acts while in the form of  
 CC two transmembrane glycoproteins, each of which is about 180 kD.  
 CC This receptor has three major regions. The first is a transmembrane  
 CC region, which spans the membrane once, separating the regions of the  
 CC receptor exterior to the cell from those interior to the cell. The  
 CC second region is an extracellular region which contains the domains  
 CC which bind the PDGF. The third region is an intracellular region  
 CC which possesses a tyrosine kinase activity. This tyrosine kinase  
 CC domain is notable in having an insert of approx. 100 amino acids,  
 CC as compared with most other receptor tyrosine kinase domains which  
 CC are contiguous or have shorter insert sequences. Fragments of this  
 CC sequence between 8 and 400 amino acids comprising one or more PDGF  
 CC ligand binding region from the extracellular domain may be used to  
 CC bind a PDGF ligand.  
 SQ Sequence 1009 AA;

Query Match 37.6%; Score 53; DB 5; Length 1009;  
 Best Local Similarity 38.1%; Pred. No. 5.69e+01;  
 Matches 8; Conservative 6; Mismatches 7; Indels 0; Gaps 0;

Db 381 vddhgstgggtvrcatgtp 401

| : : : : | : | : |

Qy 1 VIDHQGTSSKVRQKVEGS 21

## RESULT 9

ID R06910 standard; protein; 1089 AA.  
 AC R06910;  
 DT 16-JAN-1991 (first entry)  
 DE Alpha type PDGF receptor deduced from TR4 cDNA clone.  
 KW Platelet derived growth factor; Tll.

OS Homo sapiens.  
FH Key Location/Qualifiers  
FT Domain 1..23  
FT /label=signal peptide  
FT Domain 24..524  
FT /label=ligand binding domain  
FT Domain 525..548  
FT /label=transmembrane region  
FT Domain 549..599  
FT /label=juxtamembrane domain  
FT Binding-site 600..627  
FT /label=ATP binding site  
FT Modified-site 849  
FT /label=tyrosine autophosphorylation site  
FT Modified-site 42..44  
FT /label=N-glycos\_site  
FT Modified-site 76..78  
FT /label=N-glycos\_site  
FT Modified-site 103..105  
FT /label=N-glycos\_site  
FT Modified-site 179..181  
FT /label=N-glycos\_site  
FT Modified-site 353..355  
FT /label=N-glycos\_site  
FT Modified-site 359..361  
FT /label=N-glycos\_site  
FT Modified-site 458..460  
FT /label=N-glycos\_site  
FT Modified-site 468..470  
FT /label=N-glycos\_site  
FN W09010013-A.  
PD 07-SEP-1990.  
PF 08-FEB-1990; U00617.  
PR 09-FEB-1989; US-308282.  
PA (USDC ) US SEC OF COMMERCE.  
PI Matsui T, Aaronson SA, Pierce JH;  
DR WPI; 90-290306/38.  
DR N-PSDB; Q05989.  
PT Type alpha platelet-derived growth factor receptor gene - useful  
PT for transforming cells to express novel protein receptor and also  
PT susceptible to genetic engineering.  
PS Claim 7; Fig 3; 64pp; English.  
CC The TR4 clone is the largest cDNA clone related to the T11 genomic  
CC clone. Isolated from a library prepd. from human thymus DNA. The  
CC T4 cDNA clone was isolated from a M426 human embryo fibroblast  
CC cDNA library. The coding region can be introduced into the pSV2  
CC gpt vector with a simian sarcoma virus LTR as a promoter and  
CC expressed in a host. The resulting protein is a novel PDGF  
CC receptor designated type alpha (the known receptor is designated  
CC type beta). The polypeptide has a calculated molecular mass of 120  
CC kD and has all the characteristics of a membrane spanning tyrosine  
CC kinase receptor. The extracellular region comprises a hydrophobic  
CC signal peptide and a ligand binding domain which has structural  
CC homology with the PDGF-R/CSF1-R subfamily. Ten Cys residues are  
CC spaced at the same positions as in other receptors of the sub-  
CC family and eight potential N-linked glycosylation sites are also  
CC present. A hydrophobic segment spans the membrane and the cyto-  
CC plasmic region comprises a juxtamembrane region, a tyrosine kinase  
CC region split into TK1 and TK2 by a hydrophilic interkinase region  
CC and a hydrophilic C-terminal tail. The TK region includes the  
CC consensus ATP binding sequence (G-X-G-X-G...K) and a tyrosine  
CC autophosphorylation site homologous to that of pp60(v-src).  
SQ Sequence 1089 AA;

Query Match 37.6%; Score 53; DB 2; Length 1089;  
Best Local Similarity 38.1%; Pred. No. 5.69e+01;  
Matches 8; Conservative 6; Mismatches 7; Indels 0; Gaps 0;  
Db 421 vddhgstgggtvrcraetp 441  
| |||: :: || ||::  
Qy 1 VIDHQGTSKCKVRQKVEGSS 21  
RESULT 10  
ID R08267 standard; protein; 1089 AA.  
AC R08267;  
DT 07-MAR-1991 (first entry)  
DE Platelet derived growth factor (PDGF) receptor protein.  
KW Atherosclerosis; fibrotic diseases.  
OS Homo sapiens.  
PN W09014425-A.  
PD 29-NOV-1990.  
PF 21-MAY-1990; U02849.  
PR 22-MAY-1989; US-355018.  
PA (ZYMO-) ZYMOGENETICS INC.  
PI Kelly JD, Murray MJ;  
DR WPI; 90-375992/50.  
DR N-PSDB; Q06869.  
PT DNA encoding platelet-derived growth factor - used to transform  
PT cells for culturing to detect PDG agonists and antagonists  
PS Claim 1; Fig 1; 30pp; English.  
CC Gene product may be expressed from a transformed cell. It has  
CC utility in detection of PDGF agonist and antagonist analogues, binding  
CC AA, AB and BB isoforms. PDGF agonists may be used to enhance wound  
CC healing, and antagonists may be used to block the effects of PDGF  
CC eg. in treatment of atherosclerosis or fibrotic diseases.  
SQ Sequence 1089 AA;  
Query Match 37.6%; Score 53; DB 2; Length 1089;  
Best Local Similarity 38.1%; Pred. No. 5.69e+01;  
Matches 8; Conservative 6; Mismatches 7; Indels 0; Gaps 0;  
Db 421 vddhgstgggtvrcraetp 441  
| |||: :: || ||::  
Qy 1 VIDHQGTSKCKVRQKVEGSS 21  
RESULT 11  
ID W04326 standard; Protein; 1196 AA.  
AC W04326;  
DT 16-JAN-1997 (first entry)  
DE Rat petrin.  
KW Petrin; neurite outgrowth associated protein; CNS;  
KW central nervous system; myelin; protein phosphatase 2C; stroke;  
KW neurodegeneration.  
OS Rattus sp.  
FH Key Location/Qualifiers  
FT Misc\_difference 129  
FT /note= "corresponds to stop codon in DNA sequence"  
FT Misc\_difference 192  
FT /note= "corresponds to stop codon in DNA sequence"  
FT Misc\_difference 205  
FT /note= "corresponds to stop codon in DNA sequence"  
FT Misc\_difference 219  
FT /note= "corresponds to stop codon in DNA sequence"  
FT Misc\_difference 225  
FT /note= "corresponds to stop codon in DNA sequence"  
FT Misc\_difference 234

FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 243
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 269
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 285
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 312
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 319
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 344
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 358
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 378
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 386
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 455
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 473
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 473
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 494
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 555
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 593
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 602
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 609
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 621
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 724
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 736
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 739
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 786
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 841
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 924
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 934
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 1017
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 1054
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 1127
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 1147
FT	/note= "corresponds to stop codon in DNA sequence"
FT	Misc_difference 1178
FT	/note= "corresponds to stop codon in DNA sequence"
PN	W09632476-A1.
PD	17-OCT-1996.
PD	12-APR-1996; CA0214.

```

PR 13-APR-1995; US-421701.
PA (MOON ) MOUNT SINAI HOSPITAL CORP.
PI Labes M, Lozano A, Roach A, Roder J;
DR WPI; 96-477127/47.
DR N-PSDB; T38484.
PT Assay for substance that modulates response of neuronal cells - and
PT neurite growth associated protein, Petrin, useful in conditions
PT involving nerve damage resulting from traumatic injury, stroke or
PT CNS degenerative disorders
PS Claim 9; Page 57-61; 119pp; English.
CC Rat petrin (M04326) is a protein involved in modulating neurite
CC growth inhibition. The amino sequence was deduced from a cDNA
CC clone (T38484) derived from an adult rat brain cDNA library; no
CC coding sequence was indicated. Petrin is a new member of the
CC protein phosphatase 2C family, and is expressed in neurons in brain
CC tissue, partic. in the Purkinje cells of the cerebellum. Petrin,
CC and antibodies raised against it, can be used to modulate neurite
CC growth and axonal regeneration.
SQ Sequence 1196 AA;

Query Match 37.6%; Score 53; DB 19; Length 1196;
Best Local Similarity 61.5%; Pred. No. 5.62e+01;
Matches 8; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

Db 900 vpshegtsphcv 912
| | | | | | | |
| | | | | | | |
Qy 1 VIDRQGTSSKCV 13

RESULT 12
ID R33439 standard; Protein; 354 AA.
AC R33439;
DT 06-JUL-1993 (first entry)
DE Ornithine cyclodeaminase C58 from Ti plasmid pTiC58
KW mu-crystallins; drug targeting; nervous acting drugs; CNS; neural;
KW neuronal; neurotransmitter agents; neuromuscular agents; NMJ;
KW neuromuscular junctions; memory agents; Alzheimers disease;
KW CNS depressants; CNS stimulants; tranquilisers; muscle relaxants;
KW antispasmodics; analgesics; anesthetics; anticonvulsants;
KW antiepileptic agents; antianxiety agents; hallucinogens; sedatives;
KW hypnotics.
KW Agrobacterium tumefaciens
PN US7844304-A.
PD 01-JAN-1993.
PF 28-FEB-1992; 844304.
PR 28-FEB-1992; US-844304.
PA (USSH ) US DEPT HEALTH & HUMAN SERVICE.
PI Kim R, Wistow G;
DI WPI; 93-093573/11.
PT New mu-crystalline proteins - having ornithine cyclodeaminase
PT activity, used in diagnosis and treatment of disorders in
PT ornithine metabolism
PS Disclosure; Page 34; 60pp; English.
CC This sequence represents ornithine cyclodeaminase (OCD) from
CC Agrobacterium Ti plasmid pTiC58. It shows approximately 30%
CC homology with the kangaroo eye lens protein mu-crystallin.
SQ Sequence 354 AA;

Query Match 36.9%; Score 52; DB 6; Length 354;
Best Local Similarity 63.6%; Pred. No. 7.25e+01;
Matches 7; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 312 ryrvdrvegss 322
| | | | | | | |
| | | | | | | |

```



Qy 11 KCVQRKVEGSS 21

RESULT 13  
 ID R14325 standard; Protein; 589 AA.  
 AC R14325;  
 DT 17-JAN-1992 (first entry)  
 DE Rat epithelin precursor.  
 KW ET; growth regulation; inhibition; stimulation.  
 OS Rattus rattus.  
 FH Key Location/Qualifiers  
 FT Protein 1..589  
 FT /label= precursor  
 FT /note= "claim 11, page 54"  
 FT Protein 280..335  
 FT /label= EP-1  
 FT /note= "claim 12, page 54"  
 FT Protein 205..261  
 FT /label= EP-2  
 FT /note= "claim 13, page 54"  
 FT Peptide 59..114  
 FT /label= EP  
 FT /note= "claim 14, page 54"  
 FT Peptide 123..179  
 FT /label= EP  
 FT /note= "claim 15, page 54"  
 FT Peptide 362..416  
 FT /label= EP  
 FT /note= "claim 16, page 54"  
 FT Peptide 440..495  
 FT /label= EP  
 FT /note= "claim 17, page 54"  
 FT Peptide 515..570  
 FT /label= EP  
 FT /note= "claim 18, page 55"  
 PN W09115510-A.  
 PD 17-OCT-1991.  
 PF 03-APR-1991; U02321.  
 PR 03-APR-1990; US-504508.  
 PR 13-MAR-1991; US-083796.  
 PA (BRIM ) BRISTOL-MYERS SQUIB.  
 PI Shoyab M, Plozman GD;  
 DR WPI; 91-325168/44.  
 DR N-PSDB; Q14338.  
 PT New cysteine-rich growth modulating proteins, epithelins - useful  
 PT as inhibitors of neoplastic cell growth and to promote wound  
 PT healing and treat psoriasis  
 PS Disclosure; Fig 18; 97pp; English.  
 CC ET-1 is a bifunctional growth regulator, capable of stimulating  
 CC the growth of some cell types while inhibiting the growth of others.  
 CC ET-2 is functionally similar to ET-1 w.r.t. growth inhibitory  
 CC bioactivity. In contrast, however, ET-2 is apparently not capable of  
 CC eliciting the growth stimulatory activity characteristic of ET-1 and,  
 CC in fact, antagonises this ET-1 activity.  
 CC See also Q14338-40, Q14952-53, R14328-9 and R15315-20.  
 SQ Sequence 589 AA;

Query Match 36.2%; Score 51; DB 3; Length 589;  
 Best Local Similarity 46.2%; Pred. No. 9.21e+01;  
 Matches 6; Conservative 5; Mismatches 2; Indels 0; Gaps 0;  
 Db 560 hceagtkclrrk 572  
 | ::::|:| |  
 Qy 4 HQGTRSKCVQRK 16

RESULT 14  
 ID R15785 standard; Protein; 914 AA.  
 AC R15785;  
 DT 10-FEB-1992 (first entry)  
 DE B.thuringiensis toxin/AcNPV gp64 fusion protein.  
 KW chimeric; fusion protein; insecticide; AcNPV; Lepidoptera larvae;  
 KW midgut targetting; bacterial endotoxin; pFAC13.  
 OS Bacillus thuringiensis var. tenebriosis.  
 OS Autographa californica Nuclear Polyhedrosis Virus.  
 PN W09117254-A.  
 PD 14-NOV-1991.  
 PF 02-MAY-1991; U03008.  
 PR 03-MAY-1990; US-518575.  
 PA (REGC ) UNIV OF CALIFORNIA.  
 PI Sivasubramanian N, Federici A;  
 DR WPI; 91-353775/48.  
 DR N-PSDB; Q14808.  
 PT Extending host range or toxicity of insecticidal proteins - using  
 PT protein capable of binding to gut epithelium of insects  
 PS Claim 55; Fig 18; 61pp; English.  
 CC A polylinker was inserted into the XmnI restriction site at the  
 CC carboxyl terminus coding region of B.thuringiensis var. tenebriosis  
 CC (Btt) toxin. DNA encoding the gp64 viral membrane protein of AcNPV  
 CC was operably linked to the Btt toxin coding sequence via the  
 CC polylinker. The gp64 gene sequences act as midgut targetting  
 CC signals for bacterial endotoxins. Plasmid pFAC13 was one of three  
 CC different Btt/gp64 gene fusions that were constructed and its  
 CC deduced amino acid sequence is given here.  
 CC See also Q14806 and Q14807.  
 SQ Sequence 914 AA;

Query Match 36.2%; Score 51; DB 3; Length 914;  
 Best Local Similarity 45.5%; Pred. No. 9.21e+01;  
 Matches 5; Conservative 5; Mismatches 1; Indels 0; Gaps 0;  
 Db 665 kfnrcikrtve 675  
 | ::::|:| |  
 Qy 8 KSKCVQRKVE 18

RESULT 15  
 ID R15784 standard; Protein; 956 AA.  
 AC R15784;  
 DT 10-FEB-1992 (first entry)  
 DE B.thuringiensis toxin/AcNPV gp64 fusion protein.  
 KW chimeric; fusion protein; insecticide; AcNPV; Lepidoptera larvae;  
 KW midgut targetting; bacterial endotoxin; pF7.  
 OS Bacillus thuringiensis var. tenebriosis.  
 OS Autographa californica Nuclear Polyhedrosis Virus.  
 PN W09117254-A.  
 PD 14-NOV-1991.  
 PF 02-MAY-1991; U03008.  
 PR 03-MAY-1990; US-518575.  
 PA (REGC ) UNIV OF CALIFORNIA.  
 PI Sivasubramanian N, Federici A;  
 DR WPI; 91-353775/48.  
 DR N-PSDB; Q14807.  
 PT Extending host range or toxicity of insecticidal proteins - using  
 PT protein capable of binding to gut epithelium of insects  
 PS Claim 55; Fig 17; 61pp; English.  
 CC A polylinker was inserted into the XmnI restriction site at the  
 CC carboxyl terminus coding region of B.thuringiensis var. tenebriosis

Jan 28 12:16

US-08-487-283A-1.rag

15

CC (Btt) toxin. DNA encoding the gp64 viral membrane protein of hcmv  
CC was operably linked to the Btt toxin coding sequence via the  
CC polylinker. The gp64 gene sequences act as midgut targeting  
CC signals for bacterial endotoxins. Plasmid pFX7 was one of three  
CC different Btt/gp64 gene fusions that were constructed and its  
CC deduced amino acid sequence is given here.  
CC See also Q14806 and Q14808.  
SQ Sequence 956 AA;

Query Match 36.2%; Score 51; DB 3; Length 956;  
Best Local Similarity 45.3%; Pred. No. 9.21e+01;  
Matches 5; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

Db 707 kfncrlkrkve 717

|::|::|

Qy 8 KSKCVQKVE 18

Search completed: Wed Jan 28 12:09:58 1998  
Job time : 14 secs.



```
#cross-references MUID:88209511
#accession A27689
##molecule_type mRNA
##residues 412-1676 ##label WET
##cross-references GB:M18879
REFERENCE A01267
#authors Fernandez, H.N.; Hugli, T.E.
#journal J. Biol. Chem. (1978) 253:6955-6164
#title Primary structural analysis of the polypeptide portion of human C5a anaphylatoxin. Polypeptide sequence determination and assignment of the oligosaccharide attachment site in C5a.
#cross-references MUID:79005687
#accession A01267
##molecule_type protein
##residues 678-751 ##label FER
REFERENCE A01266
#authors Lundwall, A.B.; Wetsel, R.A.; Kristensen, T.; Whitehead, A.S.; Woods, D.E.; Onden, R.C.; Colten, H.R.; Tack, B.F.
#journal J. Biol. Chem. (1985) 260:2108-2112
#title Isolation and sequence analysis of a cDNA clone encoding the fifth complement component.
#cross-references MUID:95130937
#accession A01266
##molecule_type mRNA
##residues 412-854,
'STALSPRLCNCIKSGCHKLRIPGSSDSPASASQVAGITGTHHQAQT'
##cross-references GB:K02874
##note the carboxyl-terminal part of the sequence in this report appears to be derived from translation of an AU repeat sequence
REFERENCE S15121
#authors Bohnsack, J.F.; Mollison, K.W.; Buko, A.M.; Ashworth, J.C.; Hill, H.R.
#journal Biochem. J. (1991) 273:635-640
#title Group B streptococci inactivate complement component C5a by enzymic cleavage at the C-terminus.
#cross-references MUID:91144547
#contents annotation
#COMMENT Complement C5 contains two disulfide-linked chains, formed by removal of four basic residues. C5 convertase releases C5a anaphylatoxin from the amino end of the alpha chain, generating C5b (beta and alpha' chains).
#COMMENT Activation of C5 initiates the spontaneous assembly of the late complement component, C5-C9, into the membrane attack complex. C5b has a transient binding site for C6. The C5b-C6 complex is the foundation upon which the membrane attack complex is assembled.
#COMMENT C5a has potent spasmogenic and chemotactic activity.
GENETICS
#gene GDB:C5
##cross-references GDB:119734
#map_position 9q33-9q33
#CLASSIFICATION #superfamily alpha-2-macroglobulin
#KEYWORDS complement alternate pathway; complement pathway; cytolysis; glycoprotein; inflammation; membrane attack complex; plasma
FEATURE
1-18 #domain signal sequence #status predicted #label SIG
19-673, 678-1676 #product complement C5 #status predicted #label MAT\
19-673, 752-1676 #product C5b #status predicted #label C5B\
19-673 #product complement C5 and C5b beta chain #status predicted #label C5BB\
678-1676 #product complement C5 alpha chain #status predicted
```

```
#label C5A
#product C5a anaphylatoxin #status experimental #label C5T\
#product C5b alpha' chain #status predicted #label C5BA\
678-751
752-1676
567-810, 634-669,
698-724, 699-731,
711-732, 866-1527;
1101-1159,
1375-1505,
1405-1474,
1520-1525,
1532-1606,
1553-1676,
1654-1657,
741
751-752
911, 1115, 1630
#disulfide bonds #status predicted\
#binding site carbohydrate (Asn) (covalent) #status experimental\
#cleavage site Arg-Leu (C5 convertase) #status experimental\
#binding site carbohydrate (Asn) (covalent) #status predicted
SUMMARY #length 1676 #molecular-weight 188330 #checksum 3858
Query Match 100.0%; Score 12048; DB 2; Length 1676;
Best Local Similarity 100.0%; Pred. No. 0.00e+00;
Matches 1676; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db 1 mqlglclclflgktwgqetvlsapklfrvgasenivlvygyteafdatiskayp 60
|||||
Qy 1 MGLGLCLCLFLGKWTGQEQYVLSAPKFRVGSASNIQVYGYTEAFDATISKYP 60
|||||
Db 61 dkkfssyqshvhlssenkfmsailltiqkqlpgqgmpvsvyvlvskhfskskmpit 120
|||||
Qy 61 DKKFSSYSGRHLSSENKFSNSAILTIQPKQLPGQGNPVSIVYLVSVSKHFSKSRMPIT 120
|||||
Db 121 ydngflfihbtkpytpdqgskvrveinddikpkakretvltfdpegsevdmvveidhi 180
|||||
Qy 121 YDNGFLFIHDTKPYTPDQSVKRVYSLNDDLKPAKRETVLTFIDPEGSEYDMVEIDHI 180
|||||
Db 181 glisfpdfkipenrygmwtikakykedfettgtayfevkeyvlphfsveieynfigy 240
|||||
Qy 181 GIISFPDFKIPSNPRYGMWTIKAKYKEDFTTGTAYFEVKEYVLPHFVSVEIEYNFIGY 240
|||||
Db 241 knfkneitikaryfynkvvteadvitfgiredlkddqkmmqtaamtmlingiaqvt 300
|||||
Qy 241 KNFKNEITIKARYFYNKVVTADVITFGIREDLKDDQKEMMQTAMNTMLINGIAQVT 300
|||||
Db 301 fdsatavkelsysledlnnkylviavtviestgfsaeaipegikvyvspklnlvatp 360
|||||
Qy 301 FDSATAVKELSYSLEDLNKKYLYIAVTVIESTGFSAEAIPEGIKVYVSPKLNLVATP 360
|||||
Db 361 lflkpgipypikvqkdsldqlvggvpvlnaqtidvngqetsdlpdskevtrvddqvaf 420
|||||
Qy 361 LFLKPGIPYPKVKQKDSLDQLVGGVPVILNAQTIDVNGQETSDLPDSKSVTRVDDGVAF 420
|||||
Db 421 vhlpsgvtvlefsvtkdaplpeengaregyraiysslsqgylidwtdhnhkallvge 480
|||||
Qy 421 VHLPSGVTVLEFVNVTKDAPLPEENQAREGYRAIAYSSLSQSYLYIDWTDNHKALLVGE 480
|||||
Db 481 hlniivtpkspydikthynylilskgkiihfgtrekfsdasyqsinipvtqnmvpsarl 540
|||||
Qy 481 HLNIVTPKSPYDIKTHYNYLILSKGKIIHFCTREKFSDSASQSQINIPVTQNMVPSRL 540
|||||
Db 541 lvvyivtqgtaelvsdsvwlneeckcngqlqwhlspdadayspgqvtvsnmatgmdaw 600
|||||
```

Qy 541 LVYVVTGQTAELVSDSVNLIEKCGNQQLVHLSPPDADATSPGQTVSLNWTACDSDSHV 600  
Db 601 alaavdsavgyvqr gakkplervfgflekedlqcaqgqlnnanvfhlaqlftltnanad 660  
Qy 601 ALAAVDSAVYGVORGAKRPLERVFQFLEKSDLGCCAGGGLNNANVFHLAGLFTLNNAD 660  
Db 661 dsqendepckelirprtliqkkieeiaakylthsvtkccydgacvndndetceqraariel 720  
Qy 661 DSQENDEPCKEILRPRTTIQKKIEETIAAKYTHSVTKCCYDCAVCNNDNETCEQRAARISL 720  
Db 721 gprcikafteccvvaqlranishkmdqlgrlhmktilpvsbkpeirsyfpeslwewhly 780  
Qy 721 GPRCIKAFTECCVVASQJLANISHKMDQLGRLHMKTILLPVSKPEIRSYFPESLWLVHLY 780  
Db 781 prrkqlqfalpdsittweiqqiglientgicvadtkakvdkdfilemniyyavvrgeqig 840  
Qy 781 PRRKQLQFALPDSITTWEIQGIGISWTGICVADTKAKVDFKDFLENNIPYSVVRGEQIQ 840  
Db 841 lkgtvnyrtagmqfcvkmavegiictseespichggtktskcurqkvegsehlvtftv 900  
Qy 841 LKGTVNYRTSGMFCVKMSAVEGICTSESPVIDHOGIKESKVRQKVEGSSHLVTFTV 900  
Db 901 lpleighlninfetwfgkailvklrvpvgkyresvgvtldprgiygtciarrkefp 960  
Qy 901 LPLEIGHLNINFSLETWFGKAILVKLTVPEGVKRESYSGVTLDPRGIYGTISRKEFP 960  
Db 961 yripdlvpykteikrlavskglvgeilsavlsqeginlthlpkgsaeelmsavvpvfy 1020  
Qy 961 YRIPDLVYPKTEIKRILSVKGLVGEILSNVLSQEGINLTHLPKGSAEELMSVVPVY 1020  
Db 1021 vfhyletgnhwnifhsdpkleqkllkklkegmleimsyrnadysvsvkgsastwlt 1080  
Qy 1021 VFHYLETGNHWNIFHSDPKLEQKLLKLLKEGMLEIMSYRNADYSVSVKGSASTWLT 1080  
Db 1081 falrvlgqvnyveqmpnsicnslilvnyqldngsfknsqgppiklqgtlpvearen 1140  
Qy 1081 FALRVLGQVNYVEQMPNSICNLSLILVNYQLDNGSFKNSQYPIKLQGTLPVEAREN 1140  
Db 1141 elyftavfivgirkafdicplvkiikalikadnfilentlpaqstftlaisayalslqdk 1200  
Qy 1141 SLVYFTAVFVIGIRKAFDICPLVKITDIALIKADNFLENTLPAQSTFTLAISAYALSIGDK 1200  
Db 1201 thpqfrsivskrealvkgnpplzrfwkdlqkdsypntgtarmvettayalltsin 1260  
Qy 1201 THPQFRSIVSKREALVKGNPPIYRFWKDLQKDSYVPNTGTARMVETAYALLTSLN 1260  
Db 1261 lkdmynvpvklseegryggqfyetqdtinalqgleyslvkqlrlmdidvsvykhk 1320  
Qy 1261 LKDMYNVPVVKLSEEGRYGGQFYETQDTINALEGLTYSLLVKQLRLMDIDVSVYKHK 1320  
Db 1321 galhnykmtdknflgrpvevillnddlivsetgfgslatvhtvtwvhtktsseevsfyik 1380  
Qy 1321 GALHNYKMTDKNFLGRPVEVILLNDLIVSETGFGSLATVHTVTWVHTKTSSEEVCSFYIK 1380  
Db 1381 idtqdesahyrgynsdykrivacasykpreessasgshavmdislpptgisaneedlk 1440  
Qy 1381 IDTQDIEASHYRGYGNSDYKRIVACASYKPREESSSGSSHAVMDISLPPTGISANEEDLK 1440  
Db 1441 alveqvqdlftdyqkdkghvilqlnspesdflcvrfrifelvfgflspatftvveyhr 1500  
Qy 1441 ALVEGVQDQLFTDQYKDKGHVILQLNSPESDFLCVRFRIEFLVFGFLSPATFTVVEYHR 1500  
Db 1501 pdkqctmftstnikikqvcsgaackcveadcgmgqellditisaetrkqtackpeiyava 1560

Qy 1501 PKQCTWFTSTNIIKIQKCEGACKCVADCGMQEELDLTISAEIRKQTACKPEIAYA 1560  
Db 1561 ykvaitsaitenvvfkykatlldiylktgeavaekdseitfikkvictnaelvkgrqylim 1620  
Qy 1561 YKVSITSITVENVFVKYKATLLDIIYKKTGEAFAEKDSEITFIKKVICTNAELVKGRQYLIM 1620  
Db 1621 gkealqikynfeyriypdlsltwiewywpdrdtccsqcaflanldefaediflncg 1676  
Qy 1621 GKEALQIKYNFERYIYPDLSLTWIEWYWPDRDTCCSSQCAFLANLDEFEDIAFLNCG 1676  
RESULT 2  
ENTRY complement C5 precursor - mouse  
TITLE C5a anaphylatoxin; C5b  
CONTAINS  
ORGANISM #formal name Mus musculus #common name house mouse  
DATE 19-Nov-1988 #sequence\_revision 15-Oct-1994 #text\_change 16-Feb-1997  
ACCESSIONS A35530; A27538; A40429  
REFERENCE A35530  
#authors Wetzel, R.A.; Fleischer, D.T.; Haviland, D.L.  
#journal J. Biol. Chem. (1990) 265:2435-2440  
#title Deficiency of the murine fifth complement component (C5). A 2-base pair gene deletion in a 5'-exon.  
#cross-references MUID:90153853  
#accession A35530  
#molecule type mRNA  
#residues 1-215, 'L' #label WET  
#cross-references GB:J05234  
REFERENCE A27538  
#authors Wetzel, R.A.; Ogata, R.T.; Tack, B.F.  
#journal Biochemistry (1987) 26:737-743  
#title Primary structure of the fifth component of murine complement.  
#cross-references MUID:87185363  
#accession A27538  
#molecule type mRNA  
#residues 'PCL', 44-1680 #label WET2  
REFERENCE A40429  
#authors Haviland, D.L.; Haviland, J.C.; Fleischer, D.T.; Wetzel, R.A.  
#journal J. Biol. Chem. (1991) 266:11818-11825  
#title Structure of the murine fifth complement component (C5) gene. A large, highly interrupted gene with a variant donor splice site and organizational homology with the third and fourth complement component genes.  
#cross-references MUID:91268053  
#accession A40429  
#molecule type DNA  
#residues 1-15 #label HAV  
#cross-references GB:M64852  
COMMENT Complement C5 contains two disulfide-linked chains, formed by removal of four basic residues. C5 convertase releases C5a anaphylatoxin from the amino end of the alpha chain, generating C5b (beta and alpha' chains).  
COMMENT Activation of C5 initiates the spontaneous assembly of the late complement components, C5-C9, into the membrane attack complex. C5b has a transient binding site for C6. The C5b-C6 complex is the foundation upon which the membrane attack complex is assembled.  
COMMENT C5a has potent spasmogenic and chemotactic activity.  
GENETICS  
#map\_position 2  
#introns 22/3; 86/3; 140/3; 164/3; 195/2; 223/1; 253/2; 291/3; 334/1; 372/3; 434/3; 502/3; 572/3; 622/3; 667/1; 691/1; 757/1;



358 ATP







#journal	J. Biol. Chem. (1975) 250:8293-8301
#title	Human anaphylatoxin (C3a) from the third component of complement.
#cross-references	MUID:76069169
#accession	A92187
##molecule_type	protein
##residues	672-680, 'N', 682-699, 'Q', 701-748 ##label HUG
REFERENCE	
#authors	Daoudaki, M.E.; Becherer, J.D.; Lambria, J.D.
#journal	J. Immunol. (1988) 140:1577-1580
#title	A 34-amino acid peptide of the third component of complement mediates properdin binding.
#cross-references	MUID:88154452
#accession	A27603
##molecule_type	protein
##residues	1409-1563 ##label DAO
REFERENCE	
#authors	Hellman, U.; Eggertsen, G.; Engstrom, A.; Sjoquist, J.
#journal	Biochem. J. (1985) 230:353-361
#title	Amino acid sequence of the trypsin-generated C3d fragment from human complement factor C3.
#cross-references	MUID:86025442
#accession	A23435
##molecule_type	protein
##residues	1002-1012, 'E', 1014-1303 ##label HEL
##note	sequence corresponding to residues 1072-1100 was not determined but was taken from de Bruijn & Fey (reference A94605)
REFERENCE	
#authors	Poznansky, M.C.; Glissold, P.M.; Lachmann, P.J.
#journal	J. Immunol. (1989) 143:1254-1258
#title	The difference between human C3F and C3S results from a single amino acid change from an asparagine to an aspartate residue at position 1216 on the alpha-chain of the complement component, C3.
#accession	A45830
##status	not compared with conceptual translation
##molecule_type	DNA
##residues	1212-1215, 'N', 1217-1223 ##label P0Z
##note	this is the C3S allele
#accession	B45830
##status	not compared with conceptual translation
##molecule_type	DNA
##residues	1212-1223 ##label P0Z
REFERENCE	
#authors	Delmer, K.; Sottrup-Jensen, L.
#journal	FEBS Lett. (1993) 315:85-90
#title	Disulfide bridges in human complement component C3b.
#contents	annotation; disulfide bonds
COMMENT	The sequence shown is the C3 fast (C3F) allele, which is found mainly in Caucasian populations and is associated with increased incidence of certain diseases.
COMMENT	Complement C3 contains two chains, formed by removal of four residues and linked by a disulfide bond. Its activation by a convertase, which is the central reaction in both classical and alternative complement pathways, releases the C3a anaphylatoxin from the amino end of the alpha chain and generates C3b, which associates with the Bb fragment of complement factor B to form the alternative-complement-pathway C3/C5 convertase.
COMMENT	C3a anaphylatoxin is a vasoactive peptide and a mediator of inflammation.
COMMENT	C3b, with its highly reactive thiol group, binds to the surface of foreign particles and facilitates phagocytosis. It binds to complement C5 and renders it susceptible to proteolysis by the

	classical-complement-pathway C3/C5 convertase. The activity of C3b is regulated by proteolytic cleavage involving factors H and I. Its degradation products can also be biologically active. The major site of synthesis of this plasma protein is the liver.
COMMENT	
GENETICS	
#gene	GCB:C3
#cross-references	GDB:119044
#map_position	19p13.3-19p13.3
#note	contains 41 exons
CLASSIFICATION	#superfamily alpha-2-macroglobulin
KEYWORDS	acute phase; complement alternate pathway; complement pathway; glycoprotein; hydrolase; immune response; inflammation; plasma; serine proteinase; thiolester bond
FEATURE	
1-22	#domain signal sequence #status predicted #label SIG\
23-667	#product complement C3 and C3b beta chain #status predicted #label C3BB\
23-667,672-1663	#product complement C3 #status predicted #label CC3\
23-667,749-1663	#product C3b #status predicted #label C3B\
672-1663	#product complement C3 alpha chain #status predicted #label CC3A\
672-748	#product C3a anaphylatoxin #status predicted #label C3T\
749-1663	#product C3b alpha' chain #status predicted #label C3BA\
946-1303	#product C3dk fragment #status predicted #label CDK\
955-1303	#product C3dg fragment #status predicted #label CDG\
955-1001	#product C3g fragment #status predicted #label C3G\
1002-1303	#product C3d fragment #status experimental #label C3D\
1424-1457	#region propeptin binding\
85,939	#binding_site carbohydrate (Asn) (covalent) #status experimental\
559-816,627-662,693-720,694-727,707-728,873-1513,1101-1158,1358-1489,1389-1458,1506-1511,1518-1590,1537-1661,1637-1646,748-749	#disulfide_bonds #status experimental\
954-955	#cleavage_site Arg-Ser (C3 convertase) #status predicted\
1010-1013	#cleavage_site Arg-Glu (complement factor I) #status predicted\
1303-1304	#cross-link thiolester (Cys-Gln) #status experimental\
1320-1321	#cleavage_site Arg-Ser (complement factor I) #status predicted\
1617	#cleavage_site Arg-Ser (complement factor I) #status predicted\
1617	#binding_site carbohydrate (Asn) (covalent) #status predicted\
SUMMARY	#length 1663 #molecular-weight 187163 #checksum 38
Query Match	19.6%; Score 2361; DB 2; Length 1663;
Best Local Similarity	28.4%; Pred. No. 0.00e+00;
Matches	489; Conservative 458; Mismatches 660; Indels 115; Gaps 91;
Db	8 all-11-11th1pla1g-spms1i1pn1rlseetm1vleahdagqdvptvtvhdfpg 64
Qy	2 GLIGLILCFILGK7WQEQTYYISAPK1FRVGASENV1QYGYTCFADATISIKSPD 61
Db	65 k1vl1seektv1tpatbmngvft1panrekekgkrnkftvtgafgtgvve-kv1v 123
Qy	62 KRFPSYSSGHVLLSENKFNQNSALIT10P-KQLPGQGNPVSYVLEW-SKHFSKSRMP 119







```
Qy 1516 IQKVEGACACKCEADCGMQEELDLTISAETKQCTACKPEIAYAYKVSITSITVENFV 1575
Db 1549 nylmsiltvltmqtdpenngrtfvshkqcrdalslqkqdylvwglas-dl-wvtgsr 1606
Qy 1576 KYKATLLDITVYGEAVKEDSEITFIKKVCTCTNA-ELVKGROYLIMCKEALQIKYNSFR 1634
Db 1607 fyyliiskdwleapwleescqdadlqplcqdftfcsdnlvlfgc 1650
Qy 1635 YIYPLDLSLTWIEYPRDTCSSCQ-AFL-ANLDEPAEDIFLNGC 1676

RESULT 7
ENTRY C3MS #type complete
TITLE complement C3 precursor - mouse
CONTAINS alternative-complement-pathway C3/C5 convertase (EC
3.4.21.47) C3b subunit; C3a anaphylatoxin
ORGANISM #formal name Mus musculus #common name house mouse
DATE 30-Jun-1988 #sequence_revision 30-Jun-1988 #text_change
06-Sep-1996
ACCESSIONS A92459; B92459; A92460; A93938; A21898; A54561; S16369;
S16189; I49563; I49576; A01261; A05290; A29033
REFERENCE A92459
#authors Lundwall, A.; Wetzel, R.A.; Domdey, H.; Tack, B.F.; Fey, G.H.
#journal J. Biol. Chem. (1984) 259:13851-13856
#title Structure of murine complement component C3: I. Nucleotide
sequence of cloned complementary and genomic DNA coding for
the beta chain.
#cross-references MUID:85054818
#accession A92459
#molecule_type mRNA
#residues 1-724 ##label LU1
#accession B92459
#molecule_type DNA
#residues 1-124 ##label LU2
REFERENCE A92460
#authors Wetzel, R.A.; Lundwall, A.; Davidson, F.; Gibson, T.; Tack,
B.F.; Fey, G.H.
#journal J. Biol. Chem. (1984) 259:13857-13862
#title Structure of murine complement component C3: II. Nucleotide
sequence of cloned complementary DNA coding for the alpha
chain.
#cross-references MUID:85054819
#accession A92460
#molecule_type mRNA
#residues 671-1663 ##label WET
REFERENCE A93938
#authors Domdey, H.; Wiebauer, K.; Kazmaier, M.; Muller, V.; Odink,
K.; Fey, G.
#journal Proc. Natl. Acad. Sci. U.S.A. (1982) 79:7619-7623
#title Characterization of the mRNA and cloned cDNA specifying the
third component of mouse complement.
#cross-references MUID:83117730
#contents C3a
#accession A93938
#molecule_type mRNA
#residues 671-748 ##label DOM
REFERENCE A21898
#authors Sottrup-Jensen, L.; Stepanik, T.M.; Kristensen, T.; Lohblad,
P.B.; Jones, C.M.; Wierzbicki, D.M.; Magnusson, S.; Domdey,
H.; Wetzel, R.A.; Lundwall, A.; Tack, B.F.; Fey, G.H.
#journal Proc. Natl. Acad. Sci. U.S.A. (1985) 82:9-13
#title Common evolutionary origin of alpha2-macroglobulin and
complement components C3 and C4.
#cross-references MUID:85113177
```

```
#accession A21898
#molecule_type mRNA
#residues 25-1663 ##label SOT
REFERENCE A54561
#authors Hamada, J.; Cavanaugh, P.G.; Miki, K.; Nicolson, G.L.
#journal Cancer Res. (1993) 53:4418-4423
#title A paracrine migration-stimulating factor for metastatic tumor
cells secreted by mouse hepatic sinusoidal endothelial
cells: identification as complement component C3b.
#accession A54561
#molecule_type protein
#residues 25-41;749-760 ##label HAM
#experimental_source migration-stimulating factor purified from medium
conditioned by mouse hepatic sinusoidal
endothelial cells
REFERENCE S16189
#authors Sato, T.; Hong, M.H.; Jin, C.H.; Ishimi, Y.; Udagawa, N.;
Shinki, T.; Abe, E.; Suda, T.
#journal FEBS Lett. (1991) 285:21-24
#title The specific production of the third component of complement
by osteoblastic cells treated with 1-alpha,
25-dihydroxyvitamin D(3).
#cross-references MUID:91293304
#accession S16369
#molecule_type protein
#residues 25-31 ##label SAT
#accession S16189
#status preliminary
#molecule_type protein
#residues 671-677, 'X', 679-680 ##label SA2
REFERENCE I49563
#authors Fey, G.; Domdey, H.; Wiebauer, K.; Whitehead, A.S.; Odink, K.
#journal Springer Semin. Immunopathol. (1983) 6:119-147
#title Structure and expression of the C3 gene.
#cross-references MUID:84045280
#accession I49563
#status preliminary
#molecule_type mRNA
#residues 25-136, 'Q', 138-240 ##label FEY
#cross-references GB:M35659; NID:g192280; CDS_PID:g192281
REFERENCE I49576
#authors Fey, G.H.; Wiebauer, K.; Domdey, H.
#journal Ann. N. Y. Acad. Sci. (1983) 421:307-312
#title Amino acid sequences of mouse complement C3 derived from
nucleotide sequences of cloned cDNA.
#cross-references MUID:84201365
#accession I49576
#status preliminary; translated from GB/EMBL/DBJ
#molecule_type mRNA
#residues 658-761 ##label RES
#cross-references GB:M30302; NID:g192391; CDS_PID:g192392
COMMENT Complement C3 contains two chains, formed by removal of four
residues and linked by a disulfide bond. Its activation by a C3
convertase, which is the central reaction in both classical and
alternative complement pathways, releases the C3a anaphylatoxin
from the amino end of the alpha chain and generates C3b, which
associates with the Bb fragment of complement factor B to form
the alternative-complement-pathway C3/C5 convertase.
COMMENT C3a anaphylatoxin is a vasoactive peptide and a mediator of
inflammation.
COMMENT C3b, with its highly reactive thiol group, binds to the surface of
foreign particles and facilitates phagocytosis. It binds to
complement C5 and renders it susceptible to proteolysis by the
classical-complement-pathway C3/C5 convertase. The activity of
```

C3b is regulated by proteolytic cleavage involving factors H and I. Its degradation products can also be biologically active.  
The major site of synthesis of this plasma protein is the liver.

COMMENT The major site of synthesis of this plasma protein is the liver.  
GENETICS 27/2; 90/3  
#intron the list of introns may be incomplete  
#note superfamily alpha-2-macroglobulin  
CLASSIFICATION acute phase; complement alternate pathway; complement  
KEYWORDS pathway; glycoprotein; hydrolase; immune response; inflammation; plasma; serine proteinase; thiolester bond

# FEATURE

1-24 #domain signal sequence #status predicted #label SIG\  
25-666 #product complement C3 and C3b beta chain #status predicted #label C3B\  
25-666, 671-1663 #product complement C3 #status predicted #label CC3\  
25-666, 749-1663 #product C3b #status predicted #label C3B\  
671-1663 #product complement C3 alpha chain #status predicted #label CC3A\  
671-748 #product C3a anaphylatoxin #status predicted #label C3TA  
749-1663 #product C3b alpha' chain #status predicted #label C3BA  
946-1303 #product C3dk fragment #status predicted #label C3DK  
1002-1303 #product C3d fragment #status predicted #label C3D\  
1424-1457 #region properdin binding

559-816, 626-661, 693-720, 694-727, 707-728, 873-1513, 1101-1158, 1358-1489, 1389-1458, 1506-1511, 1518-1590, 1537-1661, 1637-1646 #disulfide bonds #status predicted\  
748-749 #cleavage site Arg-Ser (C3 convertase) #status predicted\  
939, 1617 #binding site carbohydrate (Asn) (covalent) #status predicted\  
1010-1013 #cross-link thiolester (Cys-Gln) #status predicted\  
1303-1304 #cleavage site Arg-Ser (complement factor I) #status predicted\  
1320-1321 #cleavage site Arg-Ser (complement factor I) #status predicted

SUMMARY #length 1663 #molecular-weight 186482 #checksum 646  
Query Match 19.1%; Score 2296; DB 2; Length 1663;  
Best Local Similarity 28.2%; Pred. No. 0.00e+00;  
Matches 479; Conservative 447; Mismatches 665; Indels 106; Gaps 84;

Db 28 yailltpnrlreseetivleahdaqqdipvtvtvqdfklrqlteektvltgashlrsv 87  
Qy 23 YVISAQIRVCASENIQVYGVTEAFDATISIKSYDPDKSYSSGHHVLSSENKQNS 82  
Db 88 sikipaskfnadkedghyvtvvanfgetvvekvamvvsfsgylfictdkttytpgstvl 147  
Qy 83 ALLTIQPKLPGCONQSVSYLVVSVKHSKSRMPITYDNGFLFIHTDPVTPDOSVK 142  
Db 148 yriftvdmnlpvok-tvllitpdgipvkrdlisemqhilpls-wnipelvmagqw 205  
Qy 143 VVVYSLNDLAPKARETIVTFID-PEGSEV--DMVEEIDHIGITSFDFKIPSNRYGMW 199  
Db 206 kirafyehapqkifsaefevkeylvpsfvrveptetf-yiddnglvsviakiylgk 264  
Qy 200 TIKAKYKEDFSFTGTAFFVEKVEYVLPHPFSVSTEPEYNYFCY-KNFKNFETITKARYFNK 258

Db 265 nv-dqtavfivg-qd--gdkkislalahlrvviedvgdavltrkvlmevtrpsnadal 320  
Qy 259 VVTEADVITFGIREDLKODKQEMQAMQNTMLINGIAQVTFDSEAVKLSVYSLEDL 318  
Db 321 vqkelysvsvtilhagedmveaeresgipivtepyqihftktpkffkmpamfdlmvftnp 380  
Qy 319 NNKYLXYANTVTESTGCFSEEAIEPGIKYVLSYKLVNIVATPLFKPGIPYIKVQVKDS 378  
Db 381 -d---gs-pas-kvl-v-vtqg-sn---akaltq-ddgvaklsaintpnsrpltitvrtk 427  
Qy 379 LDQLVGGPVVILNAQTIDVNETSDLPKSVTRVDDGVSFVILNLPSSGVTVLEFNVT 438  
Db 428 kdtlpeqrqatktmeahpystmbnennyhlhsvrsmekpgdnlvnmfhlrtddgheaki 487  
Qy 439 APDLPEENQAREGVRAYAYSLSQSYLYIDWTDNHKALLVGEHLNIIIVTPKS-p-YIDKI 496  
Db 488 ryytylvnmhkgllkagrvregqdlvlsipitpefipsfrlvaytylligasgrev 547  
Qy 497 THYNLYLSKGIHFTREKFSQASYSQINIPVTQNMWPSRLVYIYVITGEQTA-EIV 555  
Db 548 adsvvvdvdsctglvkvqdrndhlapqgqcttlriegnggarvlgvavdkgvfvlmk 607  
Qy 556 SDSWNLNIEKCGQLQVHLSPDADAYSQCVTVSNMATGMSWALAAVDSAVYGVQRG 615  
Db 608 nkltqskidvvekadigctpgsknagvfmadglafktsqglqteqradlectkpaar 667  
Qy 616 AKRPLERVQFLEKSDIGCGAGGINNANVFIHLAGLFTLTNNAADDSENDPC-KEILR 674  
Db 668 rrrsvqlmermdkagqytdkglrkcedgmrdipmryscqrarlitqgencikafide 727  
Qy 675 PRRTLQ-K-KIEETAAKHSVVKKCCYDGCAC-VNNDTECEQAARISLGRICAKATEC 731  
Db 728 cnhtklreqhrdhvlgarsleediipeedilsrshfpqswlwtieelkepeknkis 787  
Qy 732 CVWASQLRANISHKD-MQLGRLLHMK-TLPLVSKPEIRSYFPESWLWEVH-L-VP-RR--- 783  
Db 788 tkvmnifkdsittwellavslsdkgicvadpyeirvmqdfidrlrlpsvsvrveqvel 847  
Qy 784 -KQQLFALPDSLTWEIOGIGISNT-GLCVADTVKAKYFKDFLEWNPISVVRGEQQL 841  
Db 848 ravlfnyrege-el--kvr-ve-l-l-hnpafcsmataknryftqikiipkpsavpyvi 900  
Qy 842 KGTWYNTYRTSGMFCVKGMSAVEGICTSESPVIDHQTKRSKCVRO-KVESSSHLVTFV 900  
Db 901 vplkiqqavevkaavfnhfnfiedgvkktlkvyvpegmrintkvaihtldpeklqggv-qk 959  
Qy 901 LPLEIGLHNFISLETM--FKKEILWTLAVVPEGVKRESYSGV-TLDPKGI-YGTISRR 956  
Db 960 vdpv-aadlsdgvdpdtdetrlil-lqgsvpvqmaedavdgerlkhlihtpagcegmimg 1017  
Qy 957 KEEPYRPL-DLVPKTEI-KRILSVKGLLVGEILSAVLSQEGINILTHLPKSAEALMS 1014  
Db 1018 mtpvtlavhlyldtqekekfgiek-rq-ealel-lkkgytqqlafkqpsaayaafnmrpp 1074  
Qy 1015 VVPYVYFHYHLETGNHNFHSDPLIEKQKLKKLKEGMLISMSYRNADYSYVWKGGSA 1074  
Db 1075 stwtlayvkvfslaanliaidshvlgavkvlilekqkpdgvfgedpvhqemigqfr 1134  
Qy 1075 STWLTAFALRWLGQVKNKYEQNQSICNSLJLMLVENYOLDNGSFKENSQYQPIKLGQTLP 1134  
Db 1135 -nateadvltafvlialqecardiceqvnslpssginkageyieasymlqirpytvaiaq 1193  
Qy 1135 VEARENSLYLTAFTVIGIRKAFDIP-LVK-IDTALIKADNFLENTLIPAQSTFTLAIISA 1192



```

8
RESULT      C3CP      #type complete
ENTRY       complement C3 precursor - guinea pig
TITLE       alternative-complement-pathway C3/C5 convertase (EC
CONTAINS    3.4.21.47) C3b subunit; C3a anaphylatoxin
ORGANISM    #formal name Cavia porcellus #common name guinea pig
DATE        01-Feb-1992 #sequence_revision 07-Oct-1994 #text_change
            16-Feb-1997
ACCSSIONS   A37156; S03375; A20342; D20342; C20342; A31222
REFERENCE   A37156
AUTHORS     Auerbach, H.S.; Burger, R.; Dodds, A.; Colten, H.R.
JOURNAL     J. Clin. Invest. (1990) 86:96-106
TITLE       Molecular basis of complement C3 deficiency in guinea pigs
CROSS-REFERENCES MUID:90307998
ACCESSION   A37156
MOLECULE_TYPE mRNA
RESIDUES    1-1666 ##label AUE
CROSS-REFERENCES GB:M34054
REFERENCE   S03375
AUTHORS     Gerard, N.P.; Lively, M.O.; Gerard, C.
JOURNAL     Protein Seq. Data Anal. (1988) 1:473-478
TITLE       Amino acid sequence of guinea pig C3a anaphylatoxin.
CROSS-REFERENCES MUID:89113342
ACCESSION   S03375
MOLECULE_TYPE protein
RESIDUES    676-730, 'N', 732-752 ##label GER
EXPERIMENTAL source complement-activated guinea pig serum

```

```

#note
form isolated is inactive C3a anaphylatoxin and is
missing the carboxyl-terminal arginine

REFERENCE
A90479
Thomas, M.L.; Tack, B.F.
Biochemistry (1983) 22:942-947
#journal
Identification and alignment of a thiol ester site in the
#title
third component of guinea pig complement.

#cross-references M01D:83178889
#accession A20342
#molecule_type protein
#residues 676-687 ##label TH1
#accession D20342
#molecule_type protein
#residues 993-1012,1014-1017,'E',1019-1030,'Y' ##label TH2
REFERENCE
A20342
Goldberger, G.; Thomas, M.L.; Tack, B.F.; Williams, J.;
Colten, H.R.; Abraham, G.N.
J. Biol. Chem. (1981) 256:12617-12619
#journal
NH2-terminal structure and cleavage of guinea pig pro-C3, the
#title
precursor of the third complement component.

#cross-references M01D:82075767
#accession C20342
#molecule_type protein
#residues 23-38 ##label GOL
COMMENT
Complement C3 contains two chains, formed by removal of four
residues and linked by a disulfide bond. Its activation by a C3
convertase, which is the central reaction in both classical and
alternative complement pathways, releases the C3a anaphylatoxin
from the amino end of the alpha chain and generates C3b, which
associates with the Bb fragment of complement factor B to form
the alternative-complement-pathway C3/C5 convertase.
COMMENT
C3a anaphylatoxin is a vasoactive peptide and a mediator of
inflammation.
COMMENT
C3b, with its highly reactive thiol group, binds to the surface of
foreign particles and facilitates phagocytosis. It binds to
complement C5 and renders it susceptible to proteolysis by the
classical-complement-pathway C3/C5 convertase. The activity of
C3b is regulated by proteolytic cleavage involving factors H and
I. Its degradation products can also be biologically active.
COMMENT
The major site of synthesis of this plasma protein is the liver.
CLASSIFICATION
#superfamily alpha-2-macroglobulin
KEYWORDS
acute phase; complement alternate pathway; complement
pathway; glycoprotein; hydrolase; immune response;
inflammation; liver; plasma; serine proteinase; thiolester
bond

FEATURE
1-22
#domain signal sequence #status predicted #label SIG\
23-671
#product complement C3 and C3b beta chain #status
predicted #label C3B8\
23-671,676-1666
#product complement C3 #status predicted #label CC3\
23-671,754-1666
#product complement C3b #status predicted #label C3B\
676-1666
#product complement C3 alpha chain #status predicted
#label CC3A\
676-753
#product C3a anaphylatoxin #status predicted #label C3Y\
754-1666
#product complement C3b alpha' chain #status predicted
#label C3BA\
951-1308
#product C3dk fragment #status predicted #label CDK\
1007-1308
#product C3d fragment #status predicted #label C3D\
1429-1461
#region properdin binding\
557-821,630-666,
698-725,699-732,
712-733,878-1517,
1106-1163,
1363-1493,

```





Db 1582 gkerrfishikrdalhlkghylmnglss-dl--wgerpmnsyilghkdtweavepae 1639  
 QY 1595 DSEITFIKKVCTNA-ELVKGRQYLINGKEALQIKYNSFRYIYP LDSLTIWIEWPRDTT 1653

Db 1640 cqdeeqqcdqldgtftennmvfvc 1664

QY 1654 C-SS-CQFLANLDEFADIFLNGC 1676

RESULT 9  
 ENTRY C3RT #type complete  
 TITLE complement C3 precursor - rat  
 ALTERNATE\_NAMES 37K phospholipase A2 inhibitory protein  
 CONTAINS alternative-complement-pathway C3/C5 convertase (EC 3.4.21.47) C3b subunit; C3a anaphylatoxin  
 ORGANISM #formal name Rattus norvegicus #common name Norway rat  
 DATE 04-Dec-1992 #sequence\_revision 07-Oct-1994 #text\_change 16-Feb-1997  
 ACCESSIONS S15764; A54562; A01260; B35979; A35979; PNO567; PNO566; A32281; S08692  
 REFERENCE S15764  
 #authors Misumi, Y.; Sohma, M.; Ikehara, Y.  
 #journal Nucleic Acids Res. (1990) 18:2178  
 #title Nucleotide and deduced amino acid sequence of rat complement C3.  
 #cross-references MUID:90245672  
 #accession S15764  
 #molecule\_type mRNA  
 #residues 1-1663 #label MIS  
 #cross-references EMBL:X52477

REFERENCE A54562  
 #authors Sundstrom, S.A.; Komm, B.S.; Ponce-de-Leon, H.; Yi, Z.; Teuscher, C.; Lyttle, C.R.  
 #journal J. Biol. Chem. (1989) 264:16941-16947  
 #title Estrogen regulation of tissue-specific expression of complement C3.  
 #accession A54562  
 #status translation not shown  
 #molecule\_type mRNA  
 #residues 'P', 1316-1595 #label SUN  
 #cross-references GB:M29866

REFERENCE A01260  
 #authors Jacobs, J.W.; Rubin, J.S.; Hugli, T.E.; Bogardt, R.A.; Mariz, I.K.; Daniels, J.S.; Daughaday, W.H.; Bradshaw, R.A.  
 #journal Biochemistry (1978) 17:5031-5038  
 #title Purification, characterization, and amino acid sequence of rat anaphylatoxin (C3a).  
 #cross-references MUID:79062262  
 #accession A01260  
 #molecule\_type protein  
 #residues 671-703, 'K', 705-720, 'KL', 723-748 #label JAC  
 #note three disulfide bonds are present

REFERENCE A35979  
 #authors Suwa, Y.; Kudo, I.; Inaizumi, A.; Okada, M.; Kamimura, T.; Suzuki, Y.; Chang, H.W.; Hara, S.; Inoue, K.  
 #journal Proc. Natl. Acad. Sci. U.S.A. (1990) 87:2395-2399  
 #title Proteinaceous inhibitors of phospholipase A-2 purified from inflammatory sites in rats.  
 #cross-references MUID:90207203  
 #accession B35979  
 #status preliminary  
 #molecule\_type protein  
 #residues 'X', 998-1005 #label SUM  
 #accession A35979

#molecule\_type protein  
 #residues 'X', 961-962, 'P', 964-969 #label SU2  
 REFERENCE PNO566  
 #authors Nakagawa, H.; Komorita, N.  
 #journal Biochem. Biophys. Res. Commun. (1993) 194:1181-1187  
 #title Complement component C3-derived neutrophil chemotactic factors purified from exudate of rat carrageenin-induced inflammation.

#accession PNO567  
 #molecule\_type protein  
 #residues 568-592 #label NAK  
 #note amino end of a C3-derived peptide designated exudate neutrophil chemotactic factor 2, or C3-beta-c  
 #accession PNO566  
 #molecule\_type protein  
 #residues 671-687 #label NAK2  
 #note amino end of peptide designated neutrophil chemotactic factor 1 and probably identical to C3a anaphylatoxin

REFERENCE A32281  
 #authors Kuivanen, P.C.; Capulong, R.B.; Harkins, R.N.; DeSombre, E.R.  
 #journal Biochem. Biophys. Res. Commun. (1989) 158:898-905  
 #title The estrogen-responsive 110K and 74K rat uterine secretory proteins are structurally related to complement component C3.

#cross-references MUID:89149812  
 #accession A32281  
 #molecule\_type protein  
 #residues 25-41 #label KU1  
 #experimental\_source I7beta-estradiol-stimulated uterus of immature rat  
 #note the authors treat this 74K uterine secretory protein, identical as far as sequenced to complement C3, as a distinct, homologous protein rather than as an alternatively processed form or fragment  
 COMMENT Complement C3 contains two chains, formed by removal of four residues and linked by a disulfide bond. Its activation by a C3 convertase, which is the central reaction in both classical and alternative complement pathways, releases the C3a anaphylatoxin from the amino end of the alpha chain and generates C3b, which associates with the Bb fragment of complement factor B to form the alternative-complement-pathway C3/C5 convertase.  
 COMMENT C3a anaphylatoxin is a vasoactive peptide and a mediator of inflammation.

COMMENT C3b, with its highly reactive thiol group, binds to the surface of foreign particles and facilitates phagocytosis. It binds to complement C5 and renders it susceptible to proteolysis by the classical-complement-pathway C3/C5 convertase. The activity of C3b is regulated by proteolytic cleavage involving factors H and I. Its degradation products can also be biologically active.  
 COMMENT The major site of synthesis of this plasma protein is the liver.

CLASSIFICATION #superfamily alpha-2-macroglobulin  
 KEYWORDS acute phase; chemotaxis; complement alternate pathway; complement pathway; glycoprotein; hydrolase; immune response; inflammation; liver; plasma; serine proteinase; thiolester bond

FEATURE  
 1-24 #domain signal sequence #status predicted #label SIG\  
 25-666 #product complement C3 and C3b beta chain #status predicted #label C3BB\  
 25-666, 671-1663 #product complement C3 #status predicted #label CC3\  
 25-666, 749-1663 #product complement C3b #status predicted #label C3B\  
 671-1663 #product complement C3 alpha chain #status predicted #label CC3A\  
 671-748 #product C3a anaphylatoxin #status experimental #label C3T\  
 C3T\

```
749-1663 #product complement C3b alpha' chain #status predicted
#label C3bA\
946-1303 #product C3dk fragment #status predicted #label CDK\
1002-1303 #product C3d fragment #status predicted #label C3D\
1424-1457 #region properdin binding\
558-816, 626-661,
693-720, 694-727,
707-728, 873-1513,
1101-1158,
1358-1489,
1389-1458,
1506-1511,
1518-1590,
1537-1661
748-749 #disulfide_bonds #status predicted\
#cleavage_site Arg-Ser (C3 convertase) #status
predicted\
939, 1617 #binding_site carbohydrate (Aen) (covalent) #status
predicted\
1010-1013 #cross-link thiolester (Cys-Gln) #status predicted\
1303-1304 #cleavage_site Arg-Ser (complement factor I) #status
predicted\
1320-1321 * #cleavage_site Arg-Ser (complement factor I) #status
predicted
SUMMARY #length 1663 #molecular-weight 186459 #checksum 3009
Query Match 18.7%; Score 2255; DB 2; Length 1663;
Best Local Similarity 28.7%; Pred. No. 0.00e+00;
Matches 478; Conservative 422; Mismatches 665; Indels 102; Gaps 76;
Db 28 ysiitpnvlrleseeefileahdaqgdvptvtvqdfi-kkqvltsektvltgatghlnr 86
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
23 YVISAPKIFRVCASENIVIQVYGEAFATISIKSPDKFYS5SGHVLSSEKQNS 82
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 87 vfiikipaskefnadqgh-kyvtvvanfgatvvekavivsfqgyliqtcktiytpgstv 145
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 83 ALLTQP-KQLPGGQNPVSYVILEWSKUF5SKSRMPITYDNGFLFIHTDKVPYTPQSV 141
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 146 fyrlftvdmllpgkvtvvietsdpvdkrdilshngyqilps-wnipelmwqgw 204
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 142 KVRYSLANDLKPAKRETVLTFIDPESEV--DMVEEIDRIGIISFPDFKIPSNRYCWM 199
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 205 kirafeyhapkqttsaefevkylpesfevlveptekey-yhgpkglevsitarflygk 263
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 200 TIRAKYKEDFTTCTATVFEVKEVYLPHF5VSTEPEYNFYGNF-KNFEITIKARYFNK 258
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 264 nv-dgtafvifgv-qd--edkklalslrlvliedsqgeavlerkvlmdgvrpspeal 319
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 259 VTEADVITGREDIKDDQKEMWQTAQNTMLINGIAQVTFDSEAVKELSYSLIEDL 318
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 320 vqkelyvsvrtilhsgsdmaersgipvtspqihktkpfkfpampfdlmvftnp 379
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 319 NNRLYIAVVTIESTGFEAEIPGKIKVLSPPYKMLVATPFLFKPGPIPIKRVQWDS 378
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 380 -d---gs-par-rvp-v-vtqg-sd---aaqltc-ddgvaklsvntpnrrqpltitvstk 426
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 379 LDQLVGVPVILNAQTIDVNOETSLDPSKSVTRVDDGVASVFLNLP5GVTVLEFNKTD 438
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 427 kegipdarqatrmqagpystmbnsmnnylhlevervelkpgdnlnvnhlrtdaqgeaki 486
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 439 APDILPEENQAREGYRAIYAVSSLSQSYLIDWTDNHRKALLVGEHLNI-I-VTPKSPYIDKI 496
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 487 ryytylvmmkgkllkagrcvqqdlvvlslpitpefipfsvrlvayvtlligangrevv 546
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 497 THNYLLILSKKILIHGTREKFSQASVQ5INIPVTQNWVPSRLLVYIYVTEQTA-ELV 555
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
```

```
Db 547 adsvvvdvdkscvgtlvvkqdpdrnqrqapqhqtllriegnqgarvgjlvavdkgvfvlmk 606
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 556 SSVWIMNIEKCGNQLQVHLS-P-DADAYSFGQTVSLNATGMD5WALAAVDSAVYGVQR 614
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 607 knkltqskldvvekadigtcpqsgknyagvfmdaqllftkngigltqdredpecakpaa 666
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 615 GAKXP LERVQFLERKSDLGCGAGGGINANNVHFLHAGLFTLTNANADD5QENDEPC-KEIL 673
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 667 rrrrsvqlmaerrmdkagtydkglrkccedgmrdimpypscqrarllitqesclkaftmd 726
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 674 RPRRTLQ-K-KTEETAARYKHSVWKKCCYDGC-VNNDETCEQRAARISLGRICAEFE 730
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 727 cnyitklreghrrdhvlgarsdvdediipeediisrshfswlwtieelkepekngi 786
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 731 CCWASQLRANISHKD-MQLRLHMK-TLLPVSKPEIRSYFPESWLEWH-L-VP-RR-- 783
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 787 stkvmmifkdsittweilavalsdkgicvadpyeitvmqdfidrlrlysvvrneqve 846
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 784 --KQLQALPDSLTTWELQIGIGISNT-GICVADTYKAKVFKDFLENNIPYSVVRGEIQ 840
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 847 iravlfnyteqek--lkvr-vellhnpafcmatakkryyqti-e-ippkssvavpyvi 900
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 841 LKGTVTNYRTSCMQFCVKMSAVEGICTSESPVIDHQTKSKCVQRKVEGSSSHLVTFV 900
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 901 vplkigldvevkaavfnhfiadgvykklkvpegmrvnktvavrtldpehlnqggvqre 960
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 901 LPLEIGLHNFLETW--FGKELLVKTLVAVPEGVKRESYSGV-TLDPRIY-GTISRR 956
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 961 dvnaadls-dqvdpdtdestril-lqgtpvamaedavdgerlkhlivtpagcgqeqnmigm 1018
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 957 KEFFPYRIPDLVPKTEI-KRILSVKGLLVCEILSAVLSQEGINILTHLPKGSRAEALMSV 1015
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1019 tptviavhyldqteqwekf-g--lekqealelikkgytqqlafkqplsaayaafnnrpps 1075
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1016 PVVYVHYLETGWHNNIFHSDP LIEKQKLKKLKEGMLSIMSYRNADYSY5VWKGG5AS 1075
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1076 twltamwrsfsalaanliaiaqvlcgavkwliilekqpdgvfqedgvihqemigfir- 1134
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1076 TWLTAFAALVLCQVKNKYVEQONQNSICNSLLMLVENYQLDNGSFKN5QYQPKLQGTLPV 1135
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1135 ntkeadvsltafvialqearldceggvmslpgslnkageyleasylnlqpyvtaiagy 1194
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1136 EARENSLYLTATVIGIRKAFDIPC-LVK-IDTALIKADNFLLIENTLPAQSTFTLAI5AY 1193
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1195 alalmnkleepy--ltkfInt-a-k-drn--r-weepgqg----l-yn-----veatsy 1235
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1194 ALSLGDKTHPQFRSIVSALKREALVKGNPPIY8FWKONLQHKDSSVPNTCTARMVETAY 1253
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1236 allaalllkdfsvppvrwlnndervyggygstqatfmvfgalagyradvpdhkdlnmd 1295
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1254 ALLTSNLKNDINYNVPVKWLSSEQRVGGGFYSTQDTINAIEGLTE5LLVKOLR-LSMD 1312
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1296 valhlparsptvfrillwesgailrseetkqnegfaltak-qkgqgtlsvvtvyhakvg 1354
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1313 IDVSYKHKGALHNYKMT-DKNFLGRPVEVLNDDLVSTGFGSLATVHTVTVHKTSTS 1371
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1355 ttctckfdlrvtlkpapeetakkpqdskesmlidictry-lg-dvdat-ms--ildsmmt 1409
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1372 EEVCS-FYLIKIDTQDIEASHYRGYNSDYKRIIVACASYPK5SREESSGSSHAVMDISLPT 1430
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Db 1410 gfipdndllessgvdryisykemdkafenkntlilylekishseedclsfkvhqhfnv 1469
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
Qy 1431 CISNEDKALVEGVDQLFTDYQI-KD-CH--VILQINSIPSSDFLCVRFRIFELEFV 1485
| | : : : : : | : : : : : | : : : : : | : : : : : |
| : : : : : | : : : : : | : : : : : | : : : : : |
```











COMMENT	Residues 1446 or 1449 may be the carboxyl end of the alpha chain.
COMMENT	There are at least two genes coding for C4 <sub>A</sub> and C4 <sub>B</sub> . Each gene has many alleles.
GENETICS	
# gene	GDB:C4A
##cross-references	GDB:119732
#map_position	6p21.3-6p21.3
#introns	22/2; 88/3; 136/1; 179/3; 209/2; 237/1; 269/2; 304/3; 349/1; 387/3; 447/3; 508/3; 570/3; 623/3; 666/1; 691/1; 757/1; 794/2; 818/1; 864/3; 934/3; 952/1; 1052/1; 1077/2; 1129/3; 1168/3; 1226/1; 1303/3; 1359/3; 1379/3; 1411/1; 1473/2; 1503/3; 1528/3; 1563/1; 1593/1; 1626/1; 1654/1; 1698/2
CLASSIFICATION	#superfamily alpha-2-macroglobulin
KEYWORDS	acute phase; complement classical pathway; glycoprotein; hydrolase; inflammation; plasma; polymorphism; serine proteinase; thiolester bond
FEATURE	
1-19	#domain signal sequence #status predicted #label SIG\
20-675	#product complement C4 beta chain #status predicted #label BET\
20-675; 757-1446,	#product C4b #status predicted #label C4B\
1454-1744	#product complement C4 alpha chain #status predicted #label ALP\
680-1446	#product C4a anaphylatoxin #status experimental #label C4A\
680-756	#region C4b-binding protein binding\
757-845	#product C4d fragment #status experimental #label C4D\
957-1336	#product complement C4 gamma chain #status predicted #label GAM\
1454-1744	#cleavage_site Arg-Ala (complement subcomponent C1s) #status experimental\
756-757	#cross-link thiolester (Cys-Gln) #status experimental\
1010-1013	#binding_site carbohydrate (Asn) (covalent) #status experimental
1328	#length 1744 #molecular-weight 192860 #checksum 9431
SUMMARY	
Query Match	14.0%; Score 1692; DB 2; Length 1744;
Best Local Similarity	28.2%; Pred. No. 5.65e-280;
Matches	466; Conservative 419; Mismatches 620; Indels 147; Gaps 116;
Db	139 ghlfqtqpiynggrvrvrfaldqkmpst-dtitvmvshgllrvkkkevmpssi 197
Qy	124 GFLFIHTDKPYTPDQVKVRVYSLNDLKKAKRETVLTFID-PEGSEVDWVEEDHIGI 182
Db	198 fq-ddfvidpiseptkwisarfdsqlesnastqfvykviplmfvbitqpgkpyiltvp 256
Qy	193 ISFPDKIPSNPRGYMTIKAKYKEDFTTGCTAYFEKVEYVLPHFSVSIPEYNI-CYK 241
Db	257 ghldemqidiqariyigkpv-qgvayvrfgllde--dqgkttfrglesqtklvmqgshis 313
Qy	242 -NFKNFEITIKARYFNKVKVTEADVIYTFGIREDKDDQKEMQMTAMQNTMLINGIAVT 300
Db	314 lkaefqdalekinm-gitdlqglrlryvaaliiyepqgemeeaeltswfvspfeldis 372
Qy	301 FD-SE--TAVKELSYYSUDELNNKYIYIAVTVYESTGGTSEAEIPGIRKVLSPYKMLV 357
Db	373 ktkrhlyvpjafllqalvrenmspsasgi.pvkvesa-tvaspgsvpevqdiqntd-gsgq 430
Qy	358 ATPLEKPGIPYPIKQVKDSDQLVGVPVILNAQITIDVNOETSDLPDSKSVTRDDGV 417
Db	431 vsipiiipqtiselqlswsagsh-pai--arltvaapp-sg-gpgflsierpds-rprp 484

Qy	418	ASFLVNLPSGVTVLEENWKTDADPLPEENQAREGYRAIAYSSLSQSQSYLYIDWTDNKHALL	477
Db	485	vqdtlnlnlravog--gatfehyymilergqv--fmnr--pkrtlltsvsvfvdhhlape	540
Qy	478	VGELHNLVITPKSPYIDKITHYNYLLSKGKIHFTREKFSZASQSIINPVTQNMVPS	537
Db	541	fyfvaify--hgdh--p--vanalrvdvqagacegkelsvld--gakqyrnqesvkhletsd	595
Qy	538	SRLVYIVTGEQTAEIVSDSWNLTEE--KCGNQLQVHLSDPADAYSQPTVSLNMAWCH	596
Db	596	lalvalgaltdalyaagekshkplnmkgvfeamnsydlcgpgbggdsalqvfqaaglaf--	654
Qy	597	DSWVALAAVDSAVYGVQRAKQPLE--RVQFLEKSDJCGGAGGGLNNANVFHLAGLTEL	654
Db	655	sdgqwtlerkrlecpekttrtkrnvnfqkainelqgyasptakrccqddqvtirlpmmr	714
Qy	655	THNADDSQENDEFC--KE--ILRPRTL--QKTEEIAAKYKHSVVKCCYDG--ACVNNDE	709
Db	715	scqtaarvaql--crepIaccqfaeslrkksrdkgqqlgraleilIgeedideddip	773
Qy	710	TCEQRAARI--SLGPRCIFAETECVCVASQLR--A--NISHKMQ--LGRLLHMKTLPLVSKPE	764
Db	774	vrsfpenwlrwvetsvdfqlllwlpdslttweihselkktgklevatpqvlrvfref	833
Qy	765	IRSYEPESWLWEVHLVPRRKQJALFALPDSLTLTMEIGISNT--GICVADTVKAKVFDV	823
Db	834	hlhrlpmsvrrfeqlrplynyldnltcvshvsvpweqlcaggglaqqvlpvags	893
Qy	824	FLEWNPYSVVRGEQIQLKQVYVNTSGMFCVKMSAVEGICTSESPVIDHGTGKSSKC	883
Db	894	arp--vafesvptaaav--slkvwarg--sefpv--g--daveksylqiekegalhreelvy	947
Qy	884	VQRKVEGSSHLVTFVPLLEIGLWVNFSLTWFGEKLELWKTLRVWPEG--VKRES--YS	940
Db	948	lnpldhrgrtlejpnsgd--pnmpd--dqdfnsyv--rvtaespldltlgse--ga--lepqqvas	10020
Qy	941	GVTLDPRGITYGTSRRKEFPYRIDLVPKTEIKRILSVKGL--LVGEILASVLQSGCINI	999
Db	1003	llrlprgcqetmiylaptlaaeryldkteqwstlppet--kd--havdllyq--gymriqqf	1059
Qy	1000	LTHLPKSAEALMSVWPVYFPHYLETGNHNIFISDPLIEKQKLRKKEGLSITMSY	1059
Db	1060	rkadgsyaawlsrdstwtlafvklvalagevggspkqlsetsnwllsq--qqadgsfq	1118
Qy	1060	RNADYSYVWKGGSASTWLTAFALRVLGKNVYQEQNSICNSLLWLVENYQDNGSEK	1119
Db	1119	dpocvldrmqggl--v--gndetvaltafvialhlhglavfdgeageplkqrveasielek	1176
Qy	1120	ENSQYQPIKLGCTLPBARENSLYTATVIGIRK--A--F--D--C--PLV--KIDTALLIK	1172
Db	1177	sfelgeaagllqahaaatayaltit--kapvdlglvahnlnlmaaqetqdn--lywsgvt	1234
Qy	1173	NFLNLTLPQA--STTTLAISAYSALSGKTHUPGRSIV--SAKREALVKGNPPYRFXKD	1230
Db	1235	qsgsnaveptpaprnpdmpqpalwettavallhlhhegkaemadqaaawlrqgs	1294
Qy	1231	NLQHKD--S--VP--N--TCT---AR--M--VETTAVALTSL--NIKDINVPNVIRKWLSEQR	1279
Db	1295	fqqgfrstqdtviaIdalsaywiashtteerglnvrlstgrngfshqlnhrqirl	1354
Qy	1280	YGCGFYSTQDTINAIEGLTEYSLL--VKQLR--LSMDID--VS---YK--HKGALHW--YK--M	1328
Db	1355	eeelqfslgekinvkvgnsgkgtlkvlrtynvldmktntcdqlqievtkghvheyvmean	1414

Qy	1329	TDK-NF-LGRPVEVLL--ND-D-LIVSTGGG-GI-AT---VHV-TTV-VH-K-T-ST5	1371
Db	1415	edyedyeyelpakddpdpaplqptqlqlfegrrrrrrreapkvveeqesrhyvtvcilwr	1474
Qy	1372	EEVCSF-Y--L--KIDTOD-IEA-SHYRGY-GNSDYKR-----IVA--C-ASYKPS-RE	1413
Db	1475	ngkvglgmaiaadvrtllogfhalradlekltalsdryshfeteqphvlylfdsavptar-	1533
Qy	1414	ESSSG--SSHAVMDISLPTGISANEEDLKALVEGVQLFTDQIKDCHVILQINSIPSSDF	1472
Db	1534	ecvqfeavgevpvqlvqpasatlidyynperricssvfvgapksrllatlcseavcqaeg	1593
Qy	1473	LCYRFRIFELFEVGFSPATFVVEYHRPDKOCTMFYST--SNIKI--QKVCEGAACKCVEA	1530
Db	1594	kcprralergldgedgymkfacyprveygfvkviredsraafletkitqvlhf	1653
Qy	1531	DCGQMGEELDITI-SAET-RKOTACK-PEIAAYKVSITSITVENFVKYKATLLDIYKT	1587
Db	1654	tkdkaaanqmrflvrasc-rlrlpqkeylimlqdg--atydlegbpqyllidsnswle	1710
Qy	1588	GEAVAEKDSIE-TFKRKVTCTNAELVKGROYLIMGKALQIKYNEFSFRYIYPLDLSLTWIE	1646
Db	1711	empeerlcrstrgaacqahndflqeygtqgc	1742
Qy	1647	YWPDRITTC-SSCQ-AFLANLDFAEDIFLNGC	1676
RESULT	14		
ENTRY	A24558	#type complete	
TITLE	complement C4 precursor - mouse		
CONTAINS	classical-complement-pathway C3/C5 convertase (EC 3.4.21.43)		
	C4b subunit; complement C4a anaphylatoxin		
ORGANISM	#formal name Mus musculus	#common name house mouse	
DATE	31-Mar-1989	#sequence_revision 11-Nov-1994	#text_change 13-Mar-1997
ACCESSIONS	A24558; A25371; A21692; A30520; A60227; A27039; A29059;		
	A01264; B41195; 159084; 148274; 154567; 169023		
REFERENCE	A24558		
#authors	Sepich, D.S.; Noonan, D.J.; Ogata, R.T.		
#journal	Proc. Natl. Acad. Sci. U.S.A. (1985) 82:5895-5899		
#title	Complete cDNA sequence of the fourth component of murine complement.		
	#cross-references MUID:85298264		
	#accession A24558		
	#molecule_type mRNA		
	#residues 1-1738	#label SEP	
	#cross-references GB:M11729		
	#experimental source strain B10.WR		
REFERENCE	A25371		
#authors	Nonaka, M.; Nakayama, K.; Yeul, Y.D.; Takahashi, M.		
#journal	J. Biol. Chem. (1985) 260:10936-10943		
#title	Complete nucleotide and derived amino acid sequences of the fourth component of mouse complement (C4). Evolutionary aspects.		
	#cross-references MUID:85289294		
	#accession A25371		
	#molecule_type mRNA		
	#residues 1-1331	'Y', 133-326, 'E', 328-569, 'E', 571-1323, 'N', 1325-1441, 'K', 1443-1452, 'V', 1454-1738	#label NON
	#cross-references GB:M11789		
	#experimental source strain FM		
REFERENCE	A94013		
#authors	Nonaka, M.; Takahashi, M.; Natsume-Sakai, S.; Nonaka, M.; Tanaka, S.; Shimizu, A.; Honjo, T.		



```

#journal      Proc. Natl. Acad. Sci. U.S.A. (1984) 81:6822-6826
#title        Isolation of cDNA clones specifying the fourth component of
              mouse complement and its isotype, sex-limited protein.
#cross-references MUID:85038607
#accession    A21692
#molecule_type mRNA
#residues     651-719, 'G', 721-738, 'AI', 741-805 ##label NO2
#cross-references GB:M12970
#experimental_source strain FM

REFERENCE
#authors      Taillon-Miller, P.A.; Shreffler, D.C.
#journal      J. Immunol. (1988) 141:2382-2387
#title        Structural basis for the C4d.1/C4d.2 serologic allotypes of
              murine complement component C4.
#cross-references MUID:89009745
#accession    A30520
#molecule_type DNA
#residues     961-1205, 'Q', 1207-1290 ##label TAI
#cross-references GB:M23186
#experimental_source strain B10.BR

REFERENCE
#accession    A60227
#authors      Ogata, R.T.; Zepf, N.E.
#journal      Eur. J. Immunol. (1990) 20:1607-1610
#title        C4 from C4-high and C4-low mouse strains have identical
              sequences in the region corresponding to the
              isotype-specific segment of human C4.
#cross-references MUID:90353398
#accession    A60227
#molecule_type DNA
#residues     1099-1142 ##label OGA
#cross-references GB:X55493

REFERENCE
#accession    A22039
#authors      Levi-Strauss, M.; Tosi, M.; Steinmetz, M.; Klein, J.; Meo, T.
#journal      Proc. Natl. Acad. Sci. U.S.A. (1985) 82:1746-1750
#title        Multiple duplications of complement C4 gene correlate with
              H-2-controlled testosterone-independent expression of its
              sex-limited isoform, C4-Slp.
#cross-references MUID:85166208
#accession    A22039
#molecule_type mRNA
#residues     1105-1118, 'A', 1120-1189, 'T', 1191-1449 ##label LEV
#cross-references GB:K02798
#experimental_source strain B10.W7R

REFERENCE
#accession    A03753
#authors      Tosi, M.; Levi-Strauss, M.; Duponchel, C.; Meo, T.
#journal      Philos. Trans. R. Soc. Lond. (1984) 306:389-394
#title        Sequence heterogeneity of murine complementary DNA clones
              related to the C4 and C4-Slp isoforms of the fourth
              complement component.
#accession    A29059
#molecule_type mRNA
#residues     1258-1376 ##label TOS
#cross-references GB:K02798

REFERENCE
#accession    A01264
#authors      Ogata, R.T.; Shreffler, D.C.; Sepich, D.S.; Lilly, S.P.
#journal      Proc. Natl. Acad. Sci. U.S.A. (1983) 80:5061-5065
#title        cDNA clone spanning the alpha-gamma subunit junction in the
              precursor of the murine fourth complement component (C4).
#cross-references MUID:83273751
#accession    A01264
#molecule_type mRNA
#residues     1360-1400, 'S', 1402-1511 ##label OCG
#cross-references GB:K00019
#experimental_source strain B10.W7R

```

```

REFERENCE
#authors      A41195
#journal      Proc. Natl. Acad. Sci. U.S.A. (1984) 81:4908-4911
#title        Genes for murine fourth complement component (C4) and
              sex-limited protein (Slp) identified by hybridization to
              C4- and Slp-specific cDNA.
#cross-references MUID:84272739
#accession    B41195
#molecule_type mRNA
#residues     1360-1400, 'S', 1402-1511 ##label OCG
#cross-references GB:K00019
#experimental_source strain B10.W7R

REFERENCE
#accession    I59084
#authors      Nonaka, M.; Kimura, H.; Yeul, Y.D.; Yokoyama, S.; Nakayama,
              K.; Takahashi, M.
#journal      Proc. Natl. Acad. Sci. U.S.A. (1986) 83:7883-7887
#title        Identification of the 5'-flanking regulatory region
              responsible for the difference in transcriptional control
              between mouse complement C4 and Slp genes.
#cross-references MUID:87017050
#accession    I59084
#status       preliminary; translated from GB/EMBL/DBJ
#molecule_type DNA
#residues     1-21 ##label RES
#cross-references GB:M14225; NID:g199291; CDS_PID:g554211

REFERENCE
#accession    I48274
#authors      Hemenway, C.; Kalff, M.; Stavenhagen, J.; Waltheil, D.;
              Robins, D.
#journal      Nucleic Acids Res. (1986) 14:2539-2554
#title        Sequence comparison of alleles of the fourth component of
              complement (C4) and sex-limited protein (Slp).
#cross-references MUID:86176748
#accession    I48274
#status       preliminary; translated from GB/EMBL/DBJ
#molecule_type mRNA
#residues     591-603, 'M', 605-1323, 'N', 1325-1452, 'V', 1454-1585, 'Q',
              1587-1738 ##label RE2
#cross-references EMBL:X05314; NID:g50241; CDS_PID:g50242

REFERENCE
#accession    I54567
#authors      Nonaka, M.; Nakayama, K.; Yeul, Y.D.; Shimizu, A.; Takahashi,
              M.
#journal      Immunol. Rev. (1985) 87:81-99
#title        Molecular cloning and characterization of complementary and
              genomic DNA clones for mouse C4 and Slp.
#cross-references MUID:86031969
#accession    I54567
#status       preliminary; translated from GB/EMBL/DBJ
#molecule_type DNA
#residues     1-128 ##label RE3
#cross-references GB:M12968; NID:g199267; CDS_PID:g199270
#accession    I69023
#status       preliminary; translated from GB/EMBL/DBJ
#molecule_type DNA
#residues     1724-1738 ##label RE4
#cross-references GB:M12969; NID:g199268; CDS_PID:g387439

COMMENT
This protein is synthesized as a single-chain precursor and, prior
to secretion, is enzymatically cleaved to form a trimer of
nonidentical chains (alpha, beta, and gamma) which are linked by
disulfide bonds.

COMMENT
The activation of complement C4 by complement subcomponent C1s
releases the C4a anaphylatoxin from the amino end of the alpha
chain and generates C4b, which associates with the 2a fragment of
complement factor 2 to form the classical-complement-pathway C3
convertase. The C4b, C2a fragment then associates with the 3b

```

fragment of complement factor 3 to form the classical-complement-pathway C5 convertase.

COMMENT C4a anaphylatoxin is a vasoactive peptide and a mediator of inflammation.

COMMENT The activity of C4b is regulated by proteolytic cleavage involving C4b-binding protein and factor I.

## GENETICS

#introns 22/2; 86/3; 977/3; 1073/3; 1125/3; 1164/3; 1221/3

#note the list of introns is incomplete

CLASSIFICATION #superfamily alpha-2-macroglobulin

KEYWORDS acute phase; complement classical pathway; glycoprotein;

hydrolase; inflammation; plasma; polymorphism; serine

proteinase; thiolster bond

## FEATURE

1-19 #domain signal sequence #status predicted #label SIG\

20-673 #product complement C4 beta chain #status predicted

#label BET\

20-673, 754-1440,

1448-1738 #product complement C4b #status predicted #label C4B\

678-1440 #product complement C4 alpha chain #status predicted

#label ALP\

678-753 #product C4a anaphylatoxin #status predicted #label C4A\

754-843 #region C4b-binding protein binding\

953-1332 #product C4d fragment #status predicted #label C4D\

1448-1738 #product complement C4 gamma chain #status predicted

#label GAM\

224, 743, 1387 #binding site carbohydrate (Asn) (covalent) #status

predicted\

753-754 #cleavage site Arg-Asn (complement subcomponent C1e)

#status predicted\

1006-1009 #cross-link thiolester (Cys-Cln) #status predicted

SUMMARY #length 1738 #molecular-weight 192870 #checksum 4149

Query Match

Best Local Similarity 13.3%; Score 1605; DB 2; Length 1738;

Matches 449; Conservative 407; Mismatches 628; Indels 143; Gaps 111;

Db 137 ghifvtdqpiyngqrvyrfaldqkmpstdfllitvsnshglrv-lkkel-ftsts 194

Qy 124 GFLEIHTDKPVYTPDQSVKVRYSINDILKPAKRETVLTFIDPEGSEVDMEIDHIGII 183

Db 195 ifqdaftidpisepqgtwkisarfedglesnrthfevkvyvlpnfekitpwkpyilmvp 254

Qy 184 SFPD-FKIPSNRYCMTIKAKYKEDFTTGATAYFEKVEYVLPHEVSIEP-E-YNFICY 240

Db 255 snsdeldidqaryigkpv-gqvaytrfaldme--gqkrtflrletqaklvgrthie 311

Qy 241 KMFKNFEITIKARYKNKVTEDADVIYTFGIEDLKDDQKEMQATAMQNTALINGIAQVT 300

Db 312 iskdfqaaalkinigrvdeglrllyaataviespggemeeaelwrfvssafsllder 371

Qy 301 FDSEATAVKELS--YYSELDANNKYLLIYAVTVIESTGCFSEAEIPGKYIKYPLKLDVA 358

Db 372 tkrhlvpghafllqalvqmsgeasvnpvkvs-a-tlvsgdsdqvlidqstngiqq-v- 428

Qy 359 TPLFLKPGIYIP IKVQVKQSLDQVGGVPVILNAQTIDNNQETSDLPDSKSVTRVDDGVA 418

Db 429 sisfpipptvtelrllvsagsl-ypai--arlvtqa-p-psrgtqflaie-pldpsrpsv 482

Qy 419 SEVLNIPSGVTVEFNKTKDADPLEENQARECYBATAYSSLSQSYLYLDWTDNHKALLV 478

Db 483 gdtfilnlpqgibapftvymyierggimamg-rep-rktv-tavsvldhqlapsv 539

Qy 479 GEHNLIIIVTPKSPYDKITHYNYLLILSKGKLIHFCTREKFSDAYSQISINIPVTQNWPPSS 538

Db 540 yfayfyhqq-h-p--vansalliniqsdceqklqlkv-d-qakeyrnadmaklriqt-dsk 594

Qy 539 RLIIYIVITGEGTAEIUSDSVWLNIEEK-CGNQLQVHLSPDADAYSFGQTVSLNMTGMD 597

Db 595 alvalavadtalyavqgrehbkpldmaskvfevinsyvcqpggddalqvfgdaglafad 654

Qy 598 SWVALAADVSAVYQORCAKQPLE--RVFQFLEKSDLCGAGGGLNANVFLAGLTFIT 655

Db 655 gdrldqtire-dlscpkckkerkrnmvfqkavseklqgysepdkrccqdgmtklpmkrt 713

Qy 656 NANADDSOENDEPC-KEIL-RPRTL--QKKIEATAKYHSHVVKCCYDGAC-VNNDDET 710

Db 714 ceqtaarvpqqa-crepfleccckfaedlrnqtresqahlarhnmhmlqeedlideddliv 772

Qy 711 CQORARISLGRPCIKAFTECCVWASQIRANISHKDMQIGRL-H-M-K--TLIPVSKPEI 765

Db 773 rtsfpbnlwrvpevdsakliltwlpdmsmttwshqvselsksgklcvakptrvrvfrkh 832

Qy 766 RSTPESKMLWEHLWPRKQLOFALPDSLITWEIQIGISNT-GICVADTVKAKVFRDVF 824

Db 833 lhlrlpisirrfefairpvlvnylnddravsvhvtvpegclagggmmaqvtpagaa 892

Qy 825 LEMNIPYSVVRGEQIQLKGTGVYNYRTSCMQFCVKMSAVEGICTSESPVHQGTSSKCV 884

Db 893 rp-vafsvvptaaanv-plkvargv-fdl----g-davskilqiekgahreelv-yn 943

Qy 885 RQKVEGSSHLVTFVLPLEIGHNINPFLTFWFKETLVKTLRVVPEG-VKRESYSGVT 943

Db 944 ldp--l-nnigtltipqsdpmvdpdgsfslvrvtasepletmgsegalspgvvasll 1000

Qy 944 LDPGIVGTISRREFFYRIPDLVPRKTEIKRILSVKGLVGEIL-S-AVLSQEGINILIT 1001

Db 1001 rlpqqaetmiylapltlaenyldrteqsklsep-kd-havdliqk-gymziqgfrk 1057

Qy 1002 HUPGSAEALMSVWPVYVYFHYLETGNHNIFHSDPLIEKQKLKKKKEGMSIMSRYN 1061

Db 1058 ndqsfqahrdstwtatavkilsaqeqvnspekloetawllaq-qlqdgshdp 1116

Qy 1062 ADYSYVWKGSGASTMTAPALRVLGQVKNYQVQNSIGNSLMLVENYQLDNGSFKN 1121

Db 1117 cpvlhramqggl-vgs-detvaltafvialhngldvfqddakqlknrvseasitkansf 1174

Qy 1122 SQYQP IKLQGTLPVEARENSLYTFTVIGIRKA--F--DIC-PL-VKIDTALIKADNF 1174

Db 1175 lqkkaagllghaataiyaltlt-kasedllrvahnslmamaeetqeh-lywglvlg 1232

Qy 1175 LLENTLPAQ-SFTTIAISAVALSIGDKTHPQFSIV-SALKREALVKGNPPIYR---F-W 1228

Db 1233 qdkvvlrptaprepvcpqapalviettavall-hlliregkkmadkaaswlthqgsf 1291

Qy 1229 KON--LQHKDSSVP-NTGT-AR-M-VETTAYALLSINIKD-I-NYVNPVIMLSEEQRY 1280

Db 1292 hqafsrtdvtldalasywiahtteekalkvrlsemgrnglktghlhnqhkvgle 1351

Qy 1281 GGCFTYQDTINAEIGUTEY---SLIV--KQLRLSMD-ID-VSYKHKA-LBNY--K-MT 1329

Db 1352 eelkfalgstievkvegnsgktlklrtynvldmknmtccqdlgievkvtagvaywdane 1411

Qy 1330 DK-NF-LGRPVEVLJ--ND-D-L-IVST-G-FG----S--GLAT-VHVTTVVHKT-STSE 1372

Db 1412 dyedydpmaaddpvpqlqvtfegrrrrrrreapkaeeesrvqvtvcirwngk 1471

Qy 1373 EVCSEY-LKI-DTQDI--EA-SHYRGY-QN-SDYKR-I--VA----C-ASYKPS-REESS 1416



Db	1707	qmcks	1711
		—	
Qy	1652	TTCSS	1656

Search completed: Wed Jan 28 12:15:48 1998  
Job time : 99 secs.



\*\*\*\*\*  
 WAPSEPA  
 \*\*\*\*\* (TM)

Release 2.1D John F. Collins, Biocomputing Research Unit.  
 Copyright (c) 1993, 1994, 1995 University of Edinburgh, U.K.  
 Distribution rights by IntelliGenetics, Inc.

MPsrch\_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Wed Jan 28 12:16:06 1998; MasPar time 37.99 Seconds  
 935.704 Million cell updates/sec

Tabular output not generated.

Title: >US-08-487-283A-2  
 Description: (1-1676) from US08487283A.pep  
 Perfect Score: 12048  
 Sequence: 1 MGLLGILCLFLFKTQGE.....CQAFIANIDFPAEDIFLNGC 1676

Scoring table: PAM 150  
 Gap 11

Searched: 59021 seqs, 21210388 residues

Post-processing: Minimum Match 0%  
 Listing first 45 summaries

Database: swiss-prot34  
 1:part1 2:part2 3:part3 4:part4 5:part5 6:part6 7:part7  
 8:part8 9:part9 10:part10 11:part11

Statistics: Mean 58.177; Variance 98.799; scale 0.589

Pred. No. is the number of results predicted by chance to have a  
 score greater than or equal to the score of the result being printed,  
 and is derived by analysis of the total score distribution.

SUMMARIES

Result	No.	Score	Match	Length	ID	Description	Pred. No.
1	12048	100.0	1676	2	CO5_HUMAN	COMPLEMENT C5 PRECURS	0.00e+00
2	9789	81.3	1680	2	CO5_MOUSE	COMPLEMENT C5 PRECURS	0.00e+00
3	2367	19.6	1651	2	CO3_NAJANA	COMPLEMENT C3 PRECURS	0.00e+00
4	2361	19.6	1663	2	CO3_HUMAN	COMPLEMENT C3 PRECURS	0.00e+00
5	2309	19.2	1673	2	CO3_LAMJA	COMPLEMENT C3 PRECURS	0.00e+00
6	2296	19.1	1663	2	CO3_MOUSE	COMPLEMENT C3 PRECURS	0.00e+00
7	2302	19.1	1666	2	CO3_CAVPO	COMPLEMENT C3 PRECURS	0.00e+00
8	2255	18.7	1663	2	CO3_RAT	COMPLEMENT C3 PRECURS	0.00e+00
9	2106	17.5	1620	2	CO3_EPTBU	COMPLEMENT C3 (CONTAI	0.00e+00
10	2002	16.6	1640	2	CO3_ONCHY	COMPLEMENT C3-1 (CONT	0.00e+00
11	1723	14.3	1741	2	CO4_HUMAN	COMPLEMENT C4 PRECURS	0.00e+00
12	1605	13.3	1738	2	CO4_MOUSE	COMPLEMENT C4 PRECURS	0.00e+00
13	796	6.6	726	2	CO3_RABIT	COMPLEMENT C3 ALPHA C	5.45e-148

14	670	5.6	1477	1	AI13_RAT	ALPHA-1-INHIBITOR III	2.99e-118
15	661	5.5	1476	1	A2MG_MOUSE	MURINOGLOBULIN 1 PREC	3.83e-116
16	532	4.8	1474	1	A2MG_HUMAN	ALPHA-2-MACROGLOBULIN	8.97e-98
17	581	4.8	1482	8	P2P_HUMAN	PREGNANCY ZONE PROTEI	1.53e-97
18	508	4.2	323	2	CO3_XENIA	COMPLEMENT C3 (FRAGME	8.31e-81
19	507	4.2	1472	1	A2MG_RAT	ALPHA-2-MACROGLOBULIN	1.40e-80
20	502	4.2	1473	7	OVO5_CHICK	OVOSTATIN PRECURSOR (	1.92e-79
21	493	4.1	1451	1	A2MH_MOUSE	MURINOGLOBULIN 2 PREC	2.11e-77
22	399	3.3	74	2	CO5A_PIG	COMPLEMENT C5A ANAPHY	2.09e-56
23	385	3.2	74	2	CO5A_BOVIN	COMPLEMENT C5A ANAPHY	2.44e-53
24	351	2.9	76	2	CO5A_RAT	COMPLEMENT C5A ANAPHY	5.68e-46
25	171	1.4	76	2	CO4A_RAT	COMPLEMENT C4A ANAPHY	3.12e-10
26	170	1.4	77	2	CO4A_BOVIN	COMPLEMENT C4A ANAPHY	4.64e-10
27	139	1.2	77	2	CO3A_PIG	COMPLEMENT C3A ANAPHY	5.44e-05
28	121	1.0	798	10	YA73_SCHPO	HYPOTHETICAL 92.1 KD	2.39e-02
29	115	1.0	1292	8	RPOC_MYCGE	DNA-DIRECTED RNA POLY	1.57e-01
30	105	0.9	257	11	YGBI_HAEIN	HYPOTHETICAL TRANSERI	3.00e+00
31	106	0.9	270	11	YD1J_SCHPO	HYPOTHETICAL 31.5 KD	2.26e+00
32	105	0.9	328	1	ASCD_YERPS	CDP-6-DEOXY-DELTA-3,4	3.00e+00
33	110	0.9	341	11	YEJK_HAEIN	HYPOTHETICAL PROTEIN	7.10e-01
34	108	0.9	461	1	AD4B_BOVIN	STEROID HORMONE RECEP	1.27e+00
35	106	0.9	462	1	AD4B_MOUSE	STEROID HORMONE RECEP	2.26e+00
36	112	0.9	472	8	PSBC_SNNY3	PHOTOSYSTEM II 44 KD	3.91e-01
37	108	0.9	589	5	KPYK_LACDE	PYRUVATE KINASE (EC 2	1.27e+00
38	108	0.9	612	10	UNC6_CAEEL	UNC-6 PROTEIN PRECURS	1.27e+00
39	108	0.9	943	5	IROA_NEIME	IRON-REGULATED OUTER	1.27e+00
40	113	0.9	1024	10	UBA1_YEAST	UBIQUITIN-ACTIVATING	2.90e-01
41	106	0.9	1030	10	VPPI_CAEEL	PUTATIVE CLATHRIN-COA	2.26e+00
42	112	0.9	1517	4	GLTB_ECOLI	GLUTAMATE SYNTHASE (N	3.91e-01
43	107	0.9	1736	11	ZOI_HUMAN	TIGHT JUNCTION PROTEIN	1.70e+00
44	108	0.9	1805	10	Y218_MYCGE	HYPOTHETICAL PROTEIN	1.27e+00
45	106	0.9	5147	3	FAT_DROME	CADHERIN-RELATED TUMO	2.26e+00

ALIGNMENTS

RESULT	1	CO5_HUMAN	STANDARD;	PRT;	1676 AA.
ID	CO5_HUMAN				
AC	P01031;				
DT	21-JUL-1986	(REL. 01, CREATED)			
DT	01-DEC-1992	(REL. 24, LAST SEQUENCE UPDATE)			
DT	01-FEB-1996	(REL. 33, LAST ANNOTATION UPDATE)			
DE	COMPLEMENT C5 PRECURSOR (CONTAINS: C5A ANAPHYLATOXIN).				
GN	C5.				
OS	HOMO SAPIENS (HUMAN).				
OC	EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;				
OC	EUTHERIA; PRIMATES.				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE; 91079575.				
RA	HAVILAND D.L., HAVILAND J.C., FLEISCHER D.T., HUNT A., WETSEL R.A.;				
RL	J. IMMUNOL. 146:362-368 (1991).				
RN	[2]				
RP	SEQUENCE OF 412-1676 FROM N.A.				
RX	MEDLINE; 88209511.				
RA	WETSEL R.A., LEMONS R.S., LEBEAU M.M., BARNUM S.R., NOACK D.,				
RA	TACK B.F.;				
RL	BIOCHEMISTRY 27:1474-1482 (1988).				
RN	[3]				
RP	SEQUENCE OF 412-902 FROM N.A.				
RX	MEDLINE; 85130937.				
RA	LUNDWALL A.B., WETSEL R.A., KRISTENSEN T., WHITEHEAD A.S.,				
RA	WOODS D.E., OGDEN R.C., COLTEN H.R., TACK B.F.;				
RL	J. BIOL. CHEM. 260:2108-2112 (1985).				

RP SEQUENCE OF 678-751.  
RX MEDLINE; 79005687.  
RA FERNANDEZ H.N., HUGLI T.E.;  
RL J. BIOL. CHEM. 253:6955-6964 (1978).  
[5]  
RP SEQUENCE OF 678-751 FROM N.A.  
RX MEDLINE; 91144574.  
RA BOHNSACK J.F., MOLLISON K.W., BUKO A.M., ASHWORTH J.C., HILL H.R.;  
RL BIOCHEM. J. 273:635-640 (1991).  
[6]  
RP STRUCTURE BY NMR OF CSA.  
RX MEDLINE; 88309754.  
RA ZUIDERWEG E.R., MOLLISON K.W., HENKIN J., CARTER G.W.;  
RL BIOCHEMISTRY 27:3568-3580 (1988).  
[7]  
RP STRUCTURE BY NMR OF CSA.  
RX MEDLINE; 89207527.  
RA ZUIDERWEG E.R., NEITTESHEIM D.G., MOLLISON K.W., CARTER G.W.;  
RL BIOCHEMISTRY 28:172-185 (1989).  
[8]  
RP STRUCTURE BY NMR OF CSA.  
RX MEDLINE; 89274164.  
RA ZUIDERWEG E.R., FESIK S.W.;  
RL BIOCHEMISTRY 28:2387-2391 (1989).  
CC -1- FUNCTION: ACTIVATION OF C5 BY A C5 CONVERTASE INITIATES THE  
CC SPONTANEOUS ASSEMBLY OF THE LATE COMPLEMENT COMPONENTS, C5-C9,  
CC INTO THE MEMBRANE ATTACK COMPLEX. CSB HAS A TRANSIENT BINDING SITE  
CC FOR C6. THE C5B-C6 COMPLEX IS THE FOUNDATION UPON WHICH THE LYtic  
CC SUBUNIT IS ASSEMBLED.  
CC -1- SUBUNIT: C5 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 BASIC  
CC RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
CC BOND. C5 CONVERTASE ACTIVATES C5 BY CLEAVING THE ALPHA CHAIN,  
CC RELEASING C5A ANAPHYLATOXIN & GENERATING C5B (BETA CHAIN + ALPHA  
CC CHAIN).  
CC -1- SIMILARITY: TO C3, C4 AND ALPHA-2-MACROGLOBULIN.  
CC -1- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C5,  
CC C5 ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
CC INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
CC PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
CC BASOPHILIC LEUKOCYTES. C5A ALSO STIMULATES THE LOCOMOTION OF  
CC POLYMORPHONUCLEAR LEUKOCYTES (CHEMOKINESIS) AND DIRECT THEIR  
CC MIGRATION TOWARD SITES OF INFLAMMATION (CHEMOTAXIS).  
CC -1- CAUTION: REF.3 SEQUENCE DIFFERS FROM THAT SHOWN FROM POSITION 855  
CC ONWARD DUE TO THE PRESENCE OF AN ALU REPEAT.  
CC -1- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.  
DR EMBL; M57729; G179983; --.  
DR EMBL; M65134; G179692; --.  
DR PIR; S15121; S15121...  
DR HSSP; P01032; 1C5A.  
DR MIM; 120900; --.  
KW PROSITE; PS00477; ALPHA 2 MACROGLOBULIN.  
DR COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; GLYCOPROTEIN;  
KW PLASMA; MEMBRANE ATTACK COMPLEX; CYTOLYSIS; INFLAMMATORY RESPONSE;  
KW SIGNAL; POLYMORPHISM.  
FT SIGNAL 1 18 POTENTIAL.  
FT CHAIN 19 673 COMPLEMENT C5 BETA CHAIN.  
FT PROPEP 674 677  
FT CHAIN 678 1676 COMPLEMENT C5 ALPHA CHAIN.  
FT PEPTIDE 678 751 C5A ANAPHYLATOXIN.  
FT CHAIN 752 1676 C5B (ALPHA').  
FT DOMAIN 698 732 ANAPHYLATOXIN-LIKE.  
FT DISULFID 698 724

FT DISULFID 699 731  
FT DISULFID 711 732  
FT CARBOHYD 741 741  
FT CARBOHYD 911 911 POTENTIAL.  
FT CARBOHYD 1115 1115 POTENTIAL.  
FT CARBOHYD 1630 1630 POTENTIAL.  
FT VARIANT 518 518 F -> S.  
SQ SEQUENCE 1676 AA; 188331 MW; 9D5C6E59 CRC32;  
Query Match 100.0%; Score 12048; DB 2; Length 1676;  
Best Local Similarity 100.0%; Pred. No. 0.00e+00;  
Matches 1676; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Db 1 mglilgicfliflfgtwtgqetvvisapkiirvgaseniivqvygtyeafdatiskayp 60  
Qy 1 MGLLGILGFLIFLGKWTGQETVVISAPKIFRVGASENIVIVQVGYTEAFDATISIKSYP 60  
Db 61 dktfesyagshvhlssenkfmsaaitiqpkqpqqnqvayvylevskhfskskmpit 120  
Qy 61 DKTSYSSGHVHLSSEKFNQSAIITIQPKLPQQNPVSYVYLEVSKHFSKSKRMPIT 120  
Db 121 ydnofifihtdkpytpdgskvrvyslnddlkpakrtetvltfdpegsevmveidhi 180  
Qy 121 YDNQFLFIHTDKPYTPDQSVKRVYSLNDDLPKAKRETVLTFTIDPEGSEVMVEIDHI 180  
Db 181 giisfpdfkipenrygmwtikakykedfettgtayfevkeyvlpbfsvsiepeynfigy 240  
Qy 181 GIISFPDFKIPSNPRYGMWTIKAKYKEDFTTGTAYFEVKEYVLPBFSVSIEPEYNFIGY 240  
Db 241 knfkneitkaryfynkvvteadvitfgiredlkddqkmmqtamqntmlinglaqt 300  
Qy 241 KNFKNEITIKARYFNKVVTADVYITFGIREDLKDDQKEMMQTAMQNTMLINGLAQVT 300  
Db 301 fsetavkelsyysledlnmkyllylvtviestqgfseaeipdqikyvsipyklnlvatp 360  
Qy 301 FSETAVKELSYYSLEDLNKYLVIYVTVIESTQGFSEAEIPGIRYVLSPIYKINLVATP 360  
Db 361 lflkpgipypikvqkdsldqlvggvvlinagtiidvncetadldpsksvtrvddqvaf 420  
Qy 361 LFLKPGIPYPIKVQKDSLDQLVGGVVLINAGTIIDVNCETSDLDPSKSVTRVDDQVAF 420  
Db 421 vlnlpsgvtvlefnvktadpdlpeenqaregyraysslsqsylyldwtDNHkallvge 480  
Qy 421 VLNLP SGVTVLEFNKTDAPDLPENQAREGYRAYSSLSQSYLYLDWTDNHKALLVGE 480  
Db 481 hlnliivtpkpeyidkithynylilekgkiihfgtrekfsdaayqsinipvtqnmvpsrl 540  
Qy 481 HLNLIIVTPKSPYIDKITHYNYLILSKGKIIHFGTREKFSDSYQSINIPVTQNMVPSRL 540  
Db 541 lvyvlytgeqtaelvsdsvlnieekcgnqlqvhlpdadavspqgtvslmmtgmdswv 600  
Qy 541 LVYVLYTGEQTAEIVSDSVLNIEEKCGNQLQVHLPDADAVSPQGTVSLMWTGMDSWV 600  
Db 601 alaavdeavvgvqrkakplervfcgletkedgcagqglinnanvfhlagltfltnanad 660  
Qy 601 ALAAVDSAVYGVQRKAKPLERVFCGLETKEDGCAGGGLINNANVFHLAGLTFLTNANAD 660  
Db 661 daqendepckellprttlqkkieiaaaykhsvvkccydcacvnmndetceqraarisl 720  
Qy 661 DSQENDEPCKELLPRRTTLQKKEIEIAAYKHSVVKCCYDCGACVNMNDETCEQRAARISL 720  
Db 721 gprcikafteccvvaqqlraniahkdmqlgrlhmktilpvskeipseyfswlwevhlv 780  
Qy 721 GPRCIKAFTECCVVASQQLRANISHKDMQLGRLHMKTILLPVSKPEIRSEYFESWLWEVHLV 780

Db 781 prtkqlqfalsdltwiegigisntgicvadtkvfkdvflemnipyvvrgeqiq 840  
|||||  
Qy 781 PRKQQLQFALSDLTWIEGIGISNTGICVADTKVKFVDVLENNIPYSVVRGSIQ 840  
Db 841 lkgvynyrtgmgfvcvksavegictseepvldhgtkastcvrqkvegssahlvctfv 900  
|||||  
Qy 841 LKGTVNYRTGMOFCVKSAAVEGICTSESPVIDHQCTSKSRQKVEGSSSHLVFTV 900  
Db 901 lplgihmifsetfwtkeilvktlrvvpegvkresysgvtldprgiygtisrrkefp 960  
|||||  
Qy 901 LPLEIGLHNFSETFMTFKEILVKTLRVVPEGVKRESYSGVTLDPRGYGTISRKEFP 960  
Db 961 yripdlvpkteikrilskvglvgeilaevisqeginlthlpkgaeealmsvvpvyf 1020  
|||||  
Qy 961 YRIPDLVPKTEIKRILSVKGLVGEILNAVLSQEGINLTHLPKGSAAELMSVVPVY 1020  
Db 1021 vfhyletgmhwnifhsdpliekqklkkllkegmlelmsvrynadysvkwkggsastwita 1080  
|||||  
Qy 1021 VFHYLETGMHWNIFHSDPLIEKQKLKKLLKEGMLELMSVRNADYSVWMKGGASWTITA 1080  
Db 1081 falrvlgvmkyvegmqmsicnallwvnyqldngsfkenskqppiklqgtlpvearen 1140  
|||||  
Qy 1081 FALRVLGVMKYVEGMQMSICNALLWVNYQLDNGSFKENSQPPIKLQGTLPVEAREN 1140  
Db 1141 slyltaftvigirkafoicplvkdialkadnflentlpaqstfclaisayalslqdk 1200  
|||||  
Qy 1141 SLYLTAFTVIGIRKAFOICPLVKDIALKADNFLENTLPAQSTFCLAISAYALS LGDK 1200  
Db 1201 thpofrsivskrealvkgppiyrfwkhldkhdssvntgtarmvettayalltsln 1260  
|||||  
Qy 1201 THPOFRSIVSKREALVKGPPIYRFWKHLDKHDSSVNTGTARMVETTAYALLTSIN 1260  
Db 1261 lkdinvmpvklwseeqyvggfygtdtinaiegltveyslvkqlrlmdldvsvkhhk 1320  
|||||  
Qy 1261 LKDINVMPVKLWSEEQYVGGFYSTQDTINAEGLTVEYSLVVKQLRLMDLDVSVKHK 1320  
Db 1321 galhnykmtkdnflgrpvevllnddlivstfgsglatvhtvtvvhkttseevcsfvlk 1380  
|||||  
Qy 1321 GALHNYKMTKDNFLGRPVEVLLNDLIVSTFGSGLATVHTVTVVHKTSTSEEVCSFYLK 1380  
Db 1381 idtqdieahryrgyngsdykrivacaaykpsreesssgeshavmdielptgisaneedlk 1440  
|||||  
Qy 1381 IDTQDIEAHRYRGYNGSDYKRIVACAAYKPSREESSSGSHAVMDISLPTGISANEEDLK 1440  
Db 1441 alvegvdqlfcdyqikdghvnlqinspsdflcvrfrifelfevgflspatftvyeyhr 1500  
|||||  
Qy 1441 ALVEGVDDQLFCDYQIKDGHVNLQINSPSDFLCVRFRIFELFEVGFLESPATFTVYEYHR 1500  
Db 1501 pdkqctmfystenikqkvgcgaackveadcgmqgeeldltiaeetrkqtackpelaya 1560  
|||||  
Qy 1501 PDKQCTMFTYSTENIKQKVGCGAACKVEADCGMQGEELDLTIAEETRKQTACKPELAYA 1560  
Db 1561 ykveitsetvenfvkykatlldyktgeavaekdseiftikvctctnaelvkgrqylm 1620  
|||||  
Qy 1561 YKVEITSETVENFVKYKATLLDYKTGEAVALKDEIFTIKVCTCTNAELVKGRQYLM 1620  
Db 1621 gtealqikynfsfrylpldsaltwiewprdtctscsqcqlanldefaediifngc 1676  
|||||  
Qy 1621 GTEALQIKYNFSFRYLPPLDSALTWIEWPRDTCTSCSQCLANLDEFAEDIIFNGC 1676

RESULT 2

ID C05\_MOUSE STANDARD; PRT; 1680 AA.

AC P06684;  
DT 01-JAN-1988 (REL. 06, CREATED)  
DT 01-NOV-1990 (REL. 16, LAST SEQUENCE UPDATE)  
DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)  
DE COMPLEMENT C5 PRECURSOR (CONTAINS: C5A ANAPHYLATOXIN).  
CN C5.  
OS MUS MUSCULUS (MOUSE).  
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;  
OC EUHERIA; RODENTIA.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE; 90153853.  
RA WETSEL R.A., FLEISCHER D.T., HAVILAND D.L.;  
RL J. BIOL. CHEM. 265:2435-2440 (1990).  
RN [2]  
RP SEQUENCE OF 41-1680 FROM N.A.  
RX MEDLINE; 87185363.  
RA WETSEL R.A., OGATA R.T., TACK B.F.;  
RL BIOCHEMISTRY 26:737-743 (1987).  
CC -!- FUNCTION: ACTIVATION OF C5 BY A C5 CONVERTASE INITIATES THE  
CC SPONTANEOUS ASSEMBLY OF THE LATE COMPLEMENT COMPONENTS, C5-C9,  
CC INTO THE MEMBRANE ATTACK COMPLEX. C5B HAS A TRANSIENT BINDING SITE  
CC FOR C6. THE C5B-C6 COMPLEX IS THE FOUNDATION UPON WHICH THE LYtic  
CC COMPLEX IS ASSEMBLED.  
CC -!- SUBUNIT: C5 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 BASIC  
CC RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
CC BOND. C5 CONVERTASE ACTIVATES C5 BY CLEAVING THE ALPHA CHAIN,  
CC RELEASING C5A ANAPHYLATOXIN & GENERATING C5B (BETA CHAIN + ALPHA'  
CC CHAIN).  
CC -!- SIMILARITY: TO C3, C4 AND ALPHA-2-MACROGLOBULIN.  
CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C5,  
CC C5 ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
CC INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
CC PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
CC BASOPHILIC LEUKOCYTES. C5A ALSO STIMULATES THE LOCOMOTION OF  
CC POLYMORPHONUCLEAR LEUKOCYTES (CHEMOKINESIS) AND DIRECT THEIR  
CC MIGRATION TOWARD SITES OF INFLAMMATION (CHEMOTAXIS).  
CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.  
CC EMBL; M35525; G309124; -.  
DR EMBL; M35526; G309123; -.  
DR PIR; A27538; A27538.  
DR PIR; A35530; A35530.  
DR HSP; P01032; 1CSA.  
DR PROSITE; PS00477; ALPHA 2 MACROGLOBULIN.  
KW COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; GLYCOPROTEIN;  
KW PLASMA; MEMBRANE ATTACK COMPLEX; CYTOLYSIS; INFLAMMATORY RESPONSE;  
KW SIGNAL.  
FT SIGNAL 1 18  
FT CHAIN 19 1680 COMPLEMENT C5.  
FT CHAIN 19 674 COMPLEMENT C5 BETA CHAIN.  
FT PROPEP 675 678  
FT CHAIN 679 1680 COMPLEMENT C5 ALPHA CHAIN.  
FT PEPTIDE 679 755 C5A ANAPHYLATOXIN.  
FT CHAIN 756 1680 C5B (ALPHA').  
FT DOMAIN 702 736 ANAPHYLATOXIN-LIKE.  
FT DISULFID 702 728 BY SIMILARITY.  
FT DISULFID 703 735 BY SIMILARITY.  
FT DISULFID 715 736 BY SIMILARITY.  
FT CARBOHYD 427 427 POTENTIAL.  
FT CARBOHYD 915 915 POTENTIAL.  
FT CARBOHYD 1119 1119 POTENTIAL.  
FT CARBOHYD 1633 1633 POTENTIAL.  
FT VARIANT 216 216 Y -> L (IN DEFECTIVE VARIANT C5D).  
FT VARIANT 217 1680 MISSING (IN DEFECTIVE VARIANT C5D).



Seq	Sequence	1680 AA; 18887 MM; AA170448 CRC32;
	Query Match	81.3%; Score 9789; DB 2; Length 1680;
	Best Local Similarity	77.8%; Pred. No. 0.00e+00;
	Matches 1307; Conservative	218; Mismatches 149; Indels 7; Gaps
Db	1 mglwgllcllflldtwwqetvysaapkllrvsgsenvvvqhgyteafdatlkeyp	60
Qy	1 MGLLGLLCLFLFLGKTKGQETVYSAPKPRVGSSENIVIQVGYTEAFDATISIKSP	60
Db	61 dkkvtfssgvmlspenkfnaalltqlnqpvreespshvylevskhfkexkkipit	120
Qy	61 DKKFSYSGGHVLSSENKFNQNSALLTQPKQLPGCGNPVSYVLYLVSKHFSSKRWPI	120
Db	121 ynnqilfhldkpytpdqgsvkiryvsgddlkpkakretvltdfidpegsevd	180
Qy	121 YNNGELFIHTDKPVTPDQSVKRVYVSNDLDJAPAKRETVLTFIDPEGSEVDMVEEIDHI	180
Db	181 qiaifpdkipsnpgywtckanykdkftttatgafeikvylprfsvsielelrfiy	240
Qy	181 QIAIFPDKIPSNPRYGWMTIKAKYKEDFTTGTAFFEYKEVYLPHFVSIEPYNFI	240
Db	241 knfnfeitkaryfynkvpdaevyafgldredildekqmmhkatqaaklvdgvaqis	300
Qy	241 KNFNFEITIKARYFYNKVTEADVYITGIREDLKDDQEMMOTAMQNTMLINGIAQVT	300
Db	301 fdsetavkelsyns ledlnhkylyiavtvteseggfseeaiepgkvkylsepytlnlvatp	360
Qy	301 FDESETAVKELSYNSLEDLNKNKYLYIAVTVIESTGFSSEAEIPGIKVYLSPYKLNULVATP	360
Db	361 lfvkpgipfsikaqvkdsleqavggvpytlnmagtvdvmdgetsdletkrsithtdgvavf	420
Qy	361 LFLKPGPIPPKQVKVNSLDQLVGGVPVLINQAQTIQNGNETSLDPSKSVTRVDDGVASF	420
Db	421 vlnlpsnvtlwfteirtdbelpenqaskeyeavyslsqsvyiviatenykplmvqe	480
Qy	421 VLNLPSPGVTVLEFENVKTDADLPENQAREYRAIAYSLSQSQSYLYIDWTDNKKALIVGE	480
Db	481 ylnlmtvtpsepvydkithynylilekqkvigtgarekifsteyqmniipvtqmppearl	540
Qy	481 YLNIVTPKSPYIDKITHYNLYLSKGIHFGTRKFSASYSINIPVTQNPVSRLL	540
Db	541 lvyiyivtgcetaelvadavwinieekcnqlqvlhlpdepyvyvypgqvtvaldmvteadswv	600
Qy	541 LVVYIVTGEQTAEIYSDSVWNIETEEKCNQLQVHLSPDADAYSQPTVSLNMAQTGMDSWV	600
Db	601 aleavdravykvqgnakramrvqfaldekedlqcgagghghdnadvflagitftlnana	660
Qy	601 ALAAVDSAVYGVQRCAPKPLERVVQFL-EKSDLCCGAGGLNANVFLAGLFTLTWANA	659
Db	661 ddshyrdscellskzrnlhlrlkeegaakykhsvpkkccydggarvfnfyetceerva	720
Qy	660 DDSQNDPECKEILRPRTLIQ--K-KIEEIAKYKHSVWVKCCYDGCVNNDTECEQRAA	716
Db	721 rvtigpclairafneccitankirpeskhpqlqrihiktllpvmkadirsyfpeswle	780
Qy	717 RISLGPRCIAFTECCVVASQLRANISHKQMLGRLLHMKTLPLPSKPEIRSYFPESWLE	776
Db	781 lhrvprkrlqvtlpdalttweiqqigisdngicvadtllakvfkvevflennipysvvrq	840
Qy	777 VHLVPRKQLQALPDSLTTWEIQIGISNTGICVADTVKAKVFKOVFLLENNIPYSVVRG	836
Db	841 eqiqlkgtyvnymtsgtkfcvkmsavegitqsgsaashltsrpsrcvqfriegeshlv	900

Qy	837	EQIQKQCTVNTVIRTSQMOCFVMSNAVEGICITSESPVIDHGTKRSKCVQRQVEGSSSHLV	896
Ddb	901	tftllpleighhsinfletsfgkdilvktllrwpegvkresyagvildpkqginvnr	960
Qy	897	TFTWLPLEGLJNINFSLTEWFKELUWTKLRWPEGVKRESYSGVTLDPRGITGYSIRR	956
Ddb	961	kefypripidlvpttkverilavkglvgefletvskeginllhlpkgsaaelmaia	1020
Qy	957	KEFFYRIPLDLVPKTEIKRILSWKGLLVGEILSAVLSEQGINILTHLPKGSAAELMSV	1016
Ddb	1021	pvfyfhyhleaaghnwimifpdtlskrglekkikvgvsvmsrynadysymkwgaaest	1080
Qy	1017	PVFYVFHYLETGNHNIFHSFDPLEKQKKLKEGLMSLTSYRNADYSYVWKGCSAST	1076
Ddb	1081	witafalrvlgvakvkgdensicnslwlvekcqlengsfkensksgvlpklqgtlpae	1140
Qy	1077	WITAFALRVLGQVKNVQEQNQSICNSLWLWENTYQJONGSFKENSTQIQIKUQTGPVE	1136
Ddb	1141	aqetllwtafavigirkavdicptmkhtaltdkadeffllentlpkstetfllaiavays	1200
Qy	1137	ARENSLYTAFVTGIRKAFDIPCLVKIDTALIKADNFLENTLPAQSTETFLAISAYALS	1196
Ddb	1201	lqdtbprfrliivaalkreaafvkgdpipyrywrdtklrpdsvpasgtagmvetayall	1260
Qy	1197	LGDKTHPQFRTVSALKREALVKALGNPPVYRFWKONIQHKOSSVPNTGTARWVETAYALL	1256
Ddb	1261	aalkkdmnyampliikweeqryggfystqdtinaieqlteyallkqihldmdinva	1320
Qy	1257	TSJLNKIDINYVNPVKWLSQEQRYGGFYSTQDTINALEGLTEYSLWKLRLSMDIDVS	1316
Ddb	1321	yhqegdfhkyktekhfgrpvevslnddlvrstgysglatvvyktvwhkvisveefcs	1380
Qy	1317	YKHKGALHNYKWKDNFLGRPVEVLNDDLLVSTGFGSLATVHTVTVWIKTSTSEEVCS	1376
Ddb	1381	fyikidtdqieasshfr-lsdegfkriiacasykpskeestsgshavmdislpitqian	1439
Qy	1377	FYIKIDTQDIEAS-HYRGYNSDKRYIVACASYKPSHEESSGSSSHAVMDISLPTGISAN	1435
Ddb	1440	eedralvegdqlltdyqikqghvilqnsipserdfclvrfeifqvgflnpattfv	1499
Qy	1436	EEDLKALVEGDQLFTDYQIKQGHVILQNSIPSSDFLCVRFREFELFEVGLSPATFTV	1495
Ddb	1500	veyhrpdqctmaysisdtrlqkvcgaactcveadcaqlqaevdlaisadrkackcp	1559
Qy	1496	YEHYHRPDQCTMFTYSNIKIOKVCEGAACKCEADCGQMQEELDTISATRKQTACKP	1555
Ddb	1560	etayaykvrkteateenvfkyatallvrytgea--adensevtfikmsctnanlvkxk	1618
Qy	1556	EIAYAYKVSITSIVENVYKYATLLDYKTGAEVAKDSEIIFIKKVTCTINAEVLKGR	1615
Ddb	1619	qylimgtveqlkhnfsfkyiypildstwieywpdtctcpscqafvenlmmfaedflns	1678
Qy	1616	QYLMGKEALQIKTNFSFRYYIYPLDUTWIEWPDRDTCSSQCAFPLANIDEFAEDFLNG	1675
Ddb	1679	c	1679
Qy	1676	c	1676
RESULT	3		
ID	CO3	NAJNA	STANDARD; PRT; 1651 AA.
AC	Q0183;		
DT	01-JUL-1993	(REL. 26, CREATED)	
OT	01-JUL-1993	(REL. 26, LAST SEQUENCE UPDATE)	

RESULT	3		
ID	CO3 NAJNA	STANDARD;	PRT; 1651 AA.
AC	Q01833;		
DT	01-JUL-1993	(REL. 26, CREATED)	
OT	01-JUL-1993	(REL. 26, LAST SEQUENCE UPDATE)	



DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)  
DE COMPLEMENT C3 PRECURSOR (CONTAINS: C3A ANAPHYLATOXIN).  
GN C3.  
OS NAJAJA NAJA (INDIAN COBRA).  
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; REPTILIA;  
ON LEPIDOSAURIA; SERPENTES.  
RV [1]  
RW SEQUENCE FROM N.A.  
RC TISSUE=LIVER;  
RX MEDLINE; 93056528.  
RA FRITZINGER D.C., CONNELLY M., PETRELIA E.C., BREDEHORST R.,  
RL VOGEL C.W.;  
RL J. IMMUNOL. 149:3554-3562(1992).  
CC -!- FUNCTION: C3 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE  
CC COMPLEMENT SYSTEM. ITS PROCESSING BY C3 CONVERTASE IS THE CENTRAL  
CC REACTION IN BOTH CLASSICAL AND ALTERNATIVE COMPLEMENT PATHWAYS.  
CC AFTER ACTIVATION C3B CAN BIND COVALENTLY, VIA ITS REACTIVE  
CC THIOLESTER, TO CELL SURFACE CARBOHYDRATES OR IMMUNE AGGREGATES.  
CC -!- SUBUNIT: C3 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 ARG  
CC RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
CC BOND. C3 CONVERTASE ACTIVATES C3 BY CLEAVING THE ALPHA CHAIN,  
CC RELEASING C3A ANAPHYLATOXIN & GENERATING C3B (BETA CHAIN + ALPHA  
CC CHAIN).  
CC -!- SIMILARITY: TO C4, C5 AND ALPHA-2-MACROGLOBULIN.  
CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C3.  
CC C3A ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
CC INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
CC PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
CC BASOPHILIC LEUKOCYTES.  
CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.  
DR EMBL; L02365; G213373; -.  
DR PIR; A46513; A46513.  
DR HSP; P01032; IC5A.  
DR PROSITE; P500477; ALPHA 2 MACROGLOBULIN.  
KW COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; PLASMA;  
KW INFLAMMATORY RESPONSE; SIGNAL.  
FT SIGNAL 1 22  
FT CHAIN 23 1651 COMPLEMENT C3.  
FT CHAIN 23 655 BETA CHAIN.  
FT CHAIN 661 1651 ALPHA CHAIN.  
FT PEPTIDE 661 738 C3A ANAPHYLATOXIN.  
FT CHAIN 739 1651 C3B (ALPHA' CHAIN).  
FT SITE 738 739 CLEAVAGE (BY C3 CONVERTASE).  
FT DOMAIN 683 718 ANAPHYLATOXIN-LIKE.  
FT DISULFID 546 807 INTERCHAIN (BY SIMILARITY).  
FT DISULFID 615 650 BY SIMILARITY.  
FT DISULFID 683 710 BY SIMILARITY.  
FT DISULFID 684 717 BY SIMILARITY.  
FT DISULFID 697 718 BY SIMILARITY.  
FT DISULFID 863 1501 BY SIMILARITY.  
FT DISULFID 1091 1147 BY SIMILARITY.  
FT DISULFID 1346 1477 BY SIMILARITY.  
FT DISULFID 1377 1446 BY SIMILARITY.  
FT DISULFID 1494 1499 BY SIMILARITY.  
FT DISULFID 1506 1578 BY SIMILARITY.  
FT DISULFID 1525 1649 BY SIMILARITY.  
FT THIOLEST 999 1002 BY SIMILARITY.  
SQ SEQUENCE 1651 AA; 184926 MW; 2958575F CRC32;

Query Match 19.6%; Score 2367; DB 2; Length 1651;  
Best Local Similarity. 30.2%; Pred. No. 0.00c+00;  
Matches 518; Conservative 416; Mismatches 655; Indels 127; Gaps 97;

Db 4 malylyvaalligfpg-s-shgalytlitpavlrtdteeqilveahgdstpkeldifvhd 61

Qy 1 MGLLGLLCELI-FLGKTMQEQTYVISAPKIFRVGASENIVQVGYTEAFDATISIKSY 59  
Db 62 prkqkltlfgervdmqagmfvtpti-kvpakehndskqngvkvvtgpgvalekvvl 120  
Qy 60 PDKKFSYSSGHVLS-SENKFNQSAIITIQKQLPGQNPVSVYVLEVSKHFSKSRMP 118  
Db 121 leysqgfvfqtgkgytpqspvrvsvfvdhnmhmdktviveftpegivsvs-ekpm 179  
Qy 119 ITYONGLELIHTDKPVYTDQSKVRVYSINDDIKPAKRETVLTFIDEGSEVDWVEID 178  
Db 180 psqair-p-ynlpelvsfqtwkavakhepsesyayfvdyreyvlpsfeyrlpsdkfl 237  
Qy 179 HIGIISPFQFKIPNRYGMWTKAKYKREDEFTTGTAYFEVKYVLPHFVSIEPEYNF 238  
Db 238 -yidgnknfhveitarylgykkv-egvafvfgvk-i-ddakkeipsdstripiiddg 292  
Qy 239 GYKN-FNFEITIKARYFNKRVVTEADVYITFGIRELDKDDQKEMQAMQNTMLINGIA 297  
Db 293 eatlkrdt-lrs-rfdlnqlvghtlysvvtitesgdmvvtteggihivtepyqiyft 350  
Qy 298 QVTFDSEAVKELSYSLIEDLANKVLYIAVTVIESTGFSSEAEIPGKYVLSPYKLV 357  
Db 351 tkpyfkpmpyeltvvtvtnp-d--gs-paa-hvp-v-vs-ea--lh-segtt-lsdgt 397  
Qy 358 ATPLEKPGIPYKVOVKDSLDQVGVPIVILNAQTIDVNOETSDLDPSKSVTRVDGV 417  
Db 398 akllintpniqslpitvtrnhgdlpterqaksmatavtqgggsnylhvaitateik 457  
Qy 418 ASFVILNPSGVTVLEFNKVTADAPLEENQAREGYRAIAYSSLSQSYLYIDMTDNHALL 477  
Db 458 pgnlpmfnvrgnanslnqikyftylilnkikfygcrprdgqlvmtlmhitpdli 517  
Qy 478 VGEHLN-I-VTPKSPYIDKITHYNYLILSKGILHFGTREFKFSASYSQINIPVTQNW 535  
Db 518 psfrfvayqv-gnn--eivadsvvvdvtdcmgtlvkvqassrddriqkpgaaikile 574  
Qy 536 PSSRLVYVYVITGQTAELVSDVWLNIEKCGNQLQVH-LSPDADAY-SPGQTVSLNMA 593  
Db 575 qdgarvgiavdkavvylndkykiskakidwtiekedfgctagsgnlgvfedaglal 634  
Qy 594 TGMDSWVALAVDSAVYGVQKAKKPLERVFQLEKSDLCGCGAGGLNANVHLAGLTF 653  
Db 635 ttstnIntkqsaakcpqpanrrrrssvllldskaskaafqdgqltkccedgmhnpmg 694  
Qy 654 LTNANADDSQENDEPCKE-ILRPRT---IQKKIEEIAAKYKHSVVRKCCYDGCACVND 709  
Db 695 ytekrakyqegdactaaflccchyikgirdenqreselflaredfedelfgdhniier 754  
Qy 710 -TCQRAARISLSPGRCIKAFTECCVWASQLRA-NISHKDMQLCRLHMKTLTPVSRKEI-R 766  
Db 755 edfpeawlteeltgpmngiesktvpfylrdsittwellavgleptkgicvaepyei 814  
Qy 767 SYPPESLMEVH-LV--PR-R----KQLQFALPDSLTITWEIQIGISNT-GICVADTVKA 817  
Db 815 tvmkdfidrlpysvvnkneqveirailynnyadedi-y-vr---vellnypafcasteg 869  
Qy 818 KYFKDVFLEWPIPVVRGEQILKGTVYNYRTSMQFCVKMSAVEGICTSPVIDHQG 877  
Db 870 qry-r-qgfpikalsaravpfvivypleqglhdeviasvrgelasdgvrkklkvpeger 927  
Qy 878 TKSKCKVRQKVEGSSRLVTFVLPLEIQLHINFSLETW---FGKEILVKTLRVWPEGV 935  
Db 928 knvrtiieldpsvkvq-ggtqeltvianklid-dkvpdtevetrievlgdpvraqiensid 985

Qy 936 RESYSGVTLP--RGVGTISRKEFPYRIPDLVPKTEIKRSLSVKGLVGEILSAVL 993  
 Db 986 geklnhiitpogcegmimtpvati-yldatqoenlgvdrte-a-ikqimt-g 1041  
 Qy 994 QEGINILTHPKGSAAELMSVP-VFYVHYLETGHNHIFHSDFLIEKQKLKKKEG 1052  
 Db 1042 yaqmvvkhkadyaaftraeswltayvkvlamasmvkdieheicggvkvllnr 1101  
 Qy 1053 MLSIWSYRNADYSYVWKSGSASWLTAFALVLCQVQYV-EQNQNSICNSLLVENVY 1111  
 Db 1102 qpqdvfknapvhlgmllggtkgaep-eas]-tafivtalleersvckeqinildesi 1158  
 Qy 1112 QLDNGSKFNSQ-QYPIKQGTLPVEARENSLYLTAFTVIGIRKAFDIC-PIVKI-DTAL 1168  
 Db 1159 nkatdyllkkyeklqpyttaltavalaadrlnhdd-r--v--lm-aa-etgrn--r-w 1207  
 Qy 1169 IKADNFELLENTLPAQSTFTLAISAYALSLGDKTHPOF-SIVSALKREALWKNPPIYRFW 1228  
 Db 1208 -e--ey-narhn-----iegtvallahlmkkaevpvrwldidkyvggtyggtq 1257  
 Qy 1229 KQNLQKQSSVPNTGARVETAYALLTSLNADINYNVPVIRKWLSEEQRYGGGFYSTQ 1288  
 Db 1258 atvmvfgaeyeiqmthqdlndisiklperevryisindnavgartvetkinedf 1317  
 Qy 1289 DTINAEIGTEYSI-LVKQIRLSMDIDVSKUKALNNYKMDKRF-L-GRPEVILNDDL 1346  
 Db 1318 tvsas-gdqkatmtiltvnaqlredanvcnkfhldvsvenelnlkqakggkaalrki 1376  
 Qy 1347 IVSTGFGSGIATVHVTVVH-KTSTSEVCS-FYIKIDTQDIEASHYRGYSGNSDYKRIVA 1404  
 Db 1377 ctry-lg-evds-tm-tiidismaltgfpdaedkrlengvdyriekfeidnmaqgt 1431  
 Qy 1405 CASYKPSRESSGSHAVMDISLPTGISANEEDIKALVEGDQLFTDYQI-KD-GH--- 1459  
 Db 1432 wviyldkvhadeclhfkhhkfhvqgsvkvsyynldeqtkfhyhpdketgln 1491  
 Qy 1460 VILQMSIPSSDFLCVFRFIELEFVCELSPATFVYHRPDQKQTFYFSNLIK--IQ 1517  
 Db 1492 kichgnicrcaetcsllnqg-k-kidqlriqacagvdyvyktilrieekdgndiy 1549  
 Qy 1518 KVCEGAACKVEADCGQMOEELDTISATRKQFACKPEIAVAYKVSITSITVENVFKY 1577  
 Db 1550 fmdvlevikggtdrnaqakatyvsqrkcealnlkldndylinqlsdlwpmk-d-dis 1607  
 Qy 1578 KATLDIYRTGEVAEKDSEITFKRVTCTNA-ELVKRCZYLING-KEAL-QIKYNSFR 1634  
 Db 1608 ylit-kn-twierwpedecq-eef-qlcddfaq 1639  
 Qy 1635 VTYELSLTWIYWPDRITTCSSCQAFIANL-DEFAE 1669

## RESULT 4

ID C03 HUMAN STANDARD; PRT; 1663 AA.  
 AC P01024;  
 DT 21-JUL-1986 (REL. 01, CREATED)  
 DT 21-JUL-1986 (REL. 01, LAST SEQUENCE UPDATE)  
 DT 01-OCT-1996 (REL. 34, LAST ANNOTATION UPDATE)  
 DE COMPLEMENT C3 PRECURSOR (CONTAINS: C3A ANAPHYLATOXIN).  
 GN C3.  
 OS HOMO SAPIENS (HUMAN).  
 OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;  
 OC EUTHERIA; PRIMATES.  
 RN (1)

RP SEQUENCE FROM N.A.  
 RX MEDLINE; 85140166.  
 RA DE BRUIJN M.H.L., FEY G.H.;  
 RL PROC. NATL. ACAD. SCI. U.S.A. 82:708-712(1985).  
 RN [2]  
 RP SEQUENCE OF 672-748.  
 RX MEDLINE; 76069169.  
 RA HUGLI T.E.;  
 RL J. BIOL. CHEM. 250:8293-8301(1975).  
 RN [3]  
 RP SEQUENCE OF 1409-1563.  
 RX MEDLINE; 88154452.  
 RA DAUDAKI M.E., BECHERER J.D., LAMBRIS J.D.;  
 RL J. IMMUNOL. 140:1577-1580(1988).  
 RN [4]  
 RP SEQUENCE OF 988-1036.  
 RX MEDLINE; 82174534.  
 RA THOMAS M.L., JANATOVA J., GRAY W.R., TACK B.F.;  
 RL PROC. NATL. ACAD. SCI. U.S.A. 79:1054-1058(1982).  
 RN [5]  
 RP STRUCTURE BY NMR OF C3A.  
 RX MEDLINE; 88276894.  
 RA NETTESHEIM D.G., EDALJI R.P., MOLLISON K.W., GREER J., ZUIDERWEG E.R.;  
 RL PROC. NATL. ACAD. SCI. U.S.A. 85:5036-5040(1988).  
 RN [6]  
 RP MUTAGENESIS OF THIOESTER BOND REGION.  
 RX MEDLINE; 92250565.  
 RA ISAAC L., ISENMAN D.E.;  
 RL J. BIOL. CHEM. 267:10062-10069(1992).  
 RN [7]  
 RP DISULFIDE BONDS.  
 RX MEDLINE; 93106233.  
 RA DOJMER K., SOTTRUP-JENSEN L.;  
 RL FEBS LETT. 315:85-90(1993).  
 RN [8]  
 RP VARIANT C3F/S.  
 RX MEDLINE; 89309808.  
 RA POZNANSKY M.C., CLISSOLD P.M., LACHMANN P.J.;  
 RL J. IMMUNOL. 143:1254-1258(1989).  
 RN [9]  
 RP ERRATUM (RETRACTION OF ABOVE ARTICLE).  
 RX MEDLINE; 90063087.  
 RA POZNANSKY M.C., CLISSOLD P.M., LACHMANN P.J.;  
 RL J. IMMUNOL. 143:3860-3862(1989).  
 RN [10]  
 RP VARIANTS GLY-102 AND PRO-314.  
 RX MEDLINE; 91011240.  
 RA BOTTO M., YONG FONG K., SO A.K., KOCH C., WALPORT M.J.;  
 RL J. EXP. MED. 172:1011-1017(1990).  
 RN [11]  
 RP VARIANT ASN-549.  
 RX MEDLINE; 95050640.  
 RA SINGER L., WHITEHEAD W.T., AKAMA H., KATZ Y., FISHELSON Z.,  
 RA WETSEL R.A.;  
 RL J. BIOL. CHEM. 269:28494-28499(1994).  
 RN [12]  
 RP VARIANT GIN-1320.  
 RA WATANABE Y., MATSUI N., YAN K., NISHIMUKAI H., TOKUNAGA K.,  
 RA JUJI T., KOBAYASHI N., KOHSAKA T.;  
 RL MOL. IMMUNOL. 30:62-62(1993).  
 CC -!- FUNCTION: C3 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE  
 CC COMPLEMENT SYSTEM. ITS PROCESSING BY C3 CONVERTASE IS THE CENTRAL  
 CC REACTION IN BOTH CLASSICAL AND ALTERNATIVE COMPLEMENT PATHWAYS.  
 CC AFTER ACTIVATION C3B CAN BIND COVALENTLY, VIA ITS REACTIVE











Qy 616 AKRQLERFQFLEKSDGCGAGGJANNANVFLHAGLUTFLTNANADDQENDEPC-KEILR 674  
Db 668 rrtvqlmrmdkagdytdqglrkcedgmrdipmryscqrarrilitqencikafidc 727  
Qy 675 PRRTIQ-K-KIEEIAARYKHSWKKCCYDGAC-VNDETCEQRAARISLPRCIRKAFTEC 731  
Db 728 cnhtklreghrdhrlglarsleediipeediilrsfpgswlwtieelkepeknigis 787  
Qy 732 CVVASQIRANISHKD-MQJLRLHMK-TLLPYSKPEIRSPESMLWEVH-L-VP-RR--- 783  
Db 788 tkvnmifkdsittwellavsldkkgcvadpveirvmqdfidrlrlpsvvrneqvei 847  
Qy 784 -KQLOFALPDLSTTWEIQIGISWT-GICVADTVKAKVFKDVFLEMMNIPYSVVRGEQIQL 841  
Db 848 ravlfmryeqe-el--kvt-ve-l-l-hnpafcmataknryfqtikippkesvavpyvi 900  
Qy 842 KGTVYNYRTSQMQFCVRMSAVEGICTSESPVIDHQTKSKCVRQ-KVEGSSHLVFTV 900  
Db 901 vplkiqqevvkaavfnhisdvkttklvvpegminktvaihtldpeklgqgv-qk 959  
Qy 901 LPLEIGHNINFSLETW--FKEELVKTLLRVPEVCVKRESYGV-TLDPRTI-YGTISRR 956  
Db 960 vdpv-aaldsqvptdtdsetril-lqspvvgmaedavderlikhlivpagceqnmig 1017  
Qy 957 KEFPYRPL-DVLPKTEI-KRLSVKGLLVGEIILSAVISOEGINILTHLPKGSAAELMS 1014  
Db 1018 mtpvavihldtqewekfgiek-tq-ealel-ikkvttqlafkqpsayaafnnrpp 1074  
Qy 1015 VVPVFFYHLETCNHNHNIHSDPLLEKQKUKKLEKMGMLSIMSTRADYSYSVWKGCSA 1074  
Db 1075 etwltayvvkfslaanliadshvlgcgvakvllilektdpdyfcdgpyvhqemiggr 1134  
Qy 1075 STWLTAFALRVKQVNVKYNQNSICNSLLMLVENYQDNGSEKNSYQPIKLOGLP 1134  
Db 1135 -nakedveltafvliatqardicqvnslpgsinksageyieasymnlqrpytvaiaq 1193  
Qy 1135 VEARENSLYLTAFTVIGIRKAFDIP-LVK-IDTALIKADNFLLNTLPAQSTFTLAISA 1192  
Db 1194 yalalmkleepy--lqkfint-a--k-drn--r-w-e-ep-dqql-yn-----veats 1234  
Qy 1193 YALSIGKTHQFRSIVSLKREALVKGNPPYRFWKDLQHKUSSVPNTGTARWETTA 1252  
Db 1235 yallalillkdfdvppvrvlneqcyvggystqatfmvfqalagyqtdvdpdhdldm 1294  
Qy 1253 YALLTSMILKIDINYVNPVIRKWLSEQRYYGGGYSTQDTINATELGTEYSLLVQLR-LSM 1311  
Db 1295 dvafhlprsaattfllwengnllrseetkmeafeltak-gkgrgtlevvayvayhakl 1353  
Qy 1312 DIDVSYKUKGALHNYKMT-DKNFLGRPVEVLNDDLLIVSTGSGLATVHVTVVHTST 1370  
Db 1354 skvtckfdllrvaipapetakkpeeaakntfleictky-lg-dvdat-ms--ildismm 1408  
Qy 1371 SEEVCS-FYIKIDTQIEASHYRCYGNSDYKRIVACASVKPSREESSGSSHAVMDISLP 1429  
Db 1409 tgfaptdkdllelasgvdrylskymnkafenkntliiylekiehteedcltfkvhyfn 1468  
Qy 1430 TGISANEEDIKALVEGDQDLTDYQI-KD-GH---VLIQINSIPSDFLCVRFRIFELFE 1484  
Db 1469 vqlipgsvkvysynleesctrfyhpdkddgmleklchseemrcraencf-mqsgoe-k 1526  
Qy 1485 VGLSPATFTVYHRPDKQCTNFI-S-TSNIKTQKVEGAACRCVEADCGQMGEILDUT 1542  
Db 1527 inlnvldkacepgvdyvykteltnkllddfdeytmliqqviksgedevqagqqrkfi 1586

Qy 1543 ISAETRQACKPEIAYAVKVSITSITVENVFVKYKATLLDLYKTCGAVERDSEITFIK 1602  
Db 1587 hikrnnalklqkkyilmwglas-dl-wgekptseyiigkdtwvehpweaeecqdkyqk 1644  
Qy 1603 KVTCTNA-ELVKGQYLIMKCALQIKYNSFRYIYPLDLSLWIEVWPRDITC-S-SQQA 1659  
Db 1645 qeelgaftesmvvygc 1661  
Qy 1660 FLANLDEAFEDIFLNGC 1676  
RESULT 7  
ID CO3 CAVPO STANDARD; PRT; 1666 AA.  
AC P12387;  
DT 01-OCT-1989 (REL. 12, CREATED)  
DT 01-JUN-1994 (REL. 29, LAST SEQUENCE UPDATE)  
DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)  
DE COMPLEMENT C3 PRECURSOR (CONTAINS: C3A ANAPHYLATOXIN).  
GN C3.  
OS CAVIA PORCELLUS (GUINEA PIG).  
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;  
OC EUTHERIA; RODENTIA.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE; 90307998.  
RA AUERBACH H.S., BURGER R., DODDS A., COLTEN H.R.;  
RL J. CLIN. INVEST. 86:96-106(1990).  
RN [2]  
RP SEQUENCE OF 676-753.  
RX MEDLINE; 89113342.  
RA GERARD N.P., LIVELY M.O., GERARD C.;  
RL PROTEIN SEQ. DATA ANAL. 1:473-478(1988).  
RN [3]  
RP SEQUENCE OF 993-1032.  
RX MEDLINE; 83178889.  
RA THOMAS M.L., TACK B.F.;  
RL BIOCHEMISTRY 22:942-947(1983).  
CC -!- FUNCTION: C3 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE  
COMPLEMENT SYSTEM. ITS PROCESSING BY C3 CONVERTASE IS THE CENTRAL  
REACTION IN BOTH CLASSICAL AND ALTERNATIVE COMPLEMENT PATHWAYS.  
AFTER ACTIVATION C3B CAN BIND COVALENTLY, VIA ITS REACTIVE  
THIOLESTER, TO CELL SURFACE CARBOHYDRATES OR IMMUNE AGGREGATES.  
CC -!- SUBUNIT: C3 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 ARG  
RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE  
BOND. C3 CONVERTASE ACTIVATES C3 BY CLEAVING THE ALPHA CHAIN,  
RELEASING C3A ANAPHYLATOXIN & GENERATING C3B (BETA CHAIN + ALPHA  
CHAIN).  
CC -!- SIMILARITY: TO C4, C5 AND ALPHA-2-MACROGLOBULIN.  
CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C3,  
C3A ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND  
BASOPHILIC LEUKOCYTES.  
CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.  
DR EMBL; M34034; G305335; -.  
DR PIR; A37156; A37156.  
DR PIR; S03375; S03375.  
DR PIR; D20342; D20342.  
DR HSSP; P01032; IC5A.  
DR PROSITE; PS00477; ALPHA\_2\_MACROGLOBULIN.  
KW COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; PLASMA;  
INFLAMMATORY RESPONSE; GLYCOPROTEIN; SIGNAL.  
FT SIGNAL 1 29







Qy 319 NKLYIATVIESGCFSEAEIPGKYVLSPIKMLVATPLFKGPIYPKQVWDS 378

Db 380 -d---ge-par-rvp-v-vtqg-ad---aaqltg-ddovaklsvntnnrqltitvstk 426

Qy 379 LDQLVGQVPIINAQTIDVNETSLDPKSKVTRVDDGVASEFVNLPSGVTVLEFNKTD 438

Db 427 kegidparqtatqaqapstymhennnylhleversvelkpgdnlnvfnhlrldaqeaki 486

Qy 439 APDLPENQAREGYRAIYASSLSQSYLYIDWTDNHKA-LVGEHNI-I-VTPKSPYDKI 496

Db 487 rrytylvnmkgllkagrvqpgdlvlslepitpefipslrvayvyligangrevv 546

Qy 497 THYNYLILSKGKLIHFCTREKFSOASVQSNIPVTQNMWPSRLLYVYVITGEQTA-ELV 555

Db 547 adsvvvdvdkscvgtlvvkgpdrnrgpaphqtdlrliegnngarvglvavdgvyfvlnk 606

Qy 556 SDSVWNIIEEKGQVQVHSLP-DADAYSPQQTVSINMATGMSWALAAVAVSVYGVQR 614

Db 607 knltqskidvvekdiaqctpgsknyagvmdagltfktngqlqtdqredpecaakpaa 666

Qy 615 GAKXP LERVFOFLEKSDLCGAGGLNANVHLAGLFTLWNAADDSQENDEPC-KEIL 673

Db 667 rrrrsqvlmermdaqytdkgllrkccedgmrdlpmypscqrarlitgagcelkafmd 726

Qy 674 RPRRTLQ-K-KIEEIAKYKSHVWKKCYDGAC-VNDETCQRAARISLGRPCIKAFTE 730

Db 727 cnyitklrghrtdhvlarsdvdedlpeedihsrhfpeawltleelkepekngr 786

Qy 731 CCVVASQILRANISHKD-MQGLRLHWK-TLLPVSKPEIRSYFPEFWLMEVH-L-VP-RR-- 783

Db 787 stkmniflksdittellavelsdkgicvadpyeitvmqffidrlrlypsvvrneqve 846

Qy 784 --KQIAQFALPDLTWEIOGIGISNT-GICVADTVKAKVDFLEWNPYSVVRGSGIQ 840

Db 847 iravlnfyreqek---lkr-vellhnpafcmatakkryqtie-ippkssvavpyvi 900

Qy 841 LKGTVYNYRSGMFCVKAASAVEGICTSESPVIDHQCKSKVRQKVGSSSHLVITV 900

Db 901 vplkiglqevvkaavnfhfiedvdkllkvpegmrvnktvavrtdlpehlnqgvqre 960

Qy 901 LPLEIGLHNFISLETW--FGKEILVKTLRVPEVKRESYSGV-TLDPRIY-GTISR 956

Db 961 dmaadls-dqvdpdsetril-lqgtvpaqmaedavdgerlklhltvpsgcqeqnimg 1018

Qy 957 KEFFYRIPDLVPKTEI-KRILSVKGLVGEILSAVLSQEGINILTHLPKGSAAELMSV 1015

Db 1019 tptviavnyldtqeakf-g--lekrqealelikkqytlqqlafkqplaaayafnrrpps 1075

Qy 1016 VPFYVYHLYETGNHNFHSDPLIEKQKLRKKKKEGMLSTMSNRADYSYVMKGGAS 1075

Db 1076 twltamwrsfelaanliaidsqvlqavkwliilektpdgvfqedpvhqemiggr- 1134

Qy 1076 TWLTFALFALVQNVYEQNSICNSLWLVENYQLDNGSFKENSQYQPIKLGQTLPV 1135

Db 1135 ntkeadvsltafvialqeadicagvmslpgelinkageyleasylnlqrptvvaigy 1194

Qy 1136 EARENSILYTAFTYVIGIRKAFDIP-LVK-IDTALIKADNFLENTLPAGSTFLAISAY 1193

Db 1195 alalmnkleepy--lktflint-a--k-drn--r-weepqgq----l-yn-----veatsy 1235

Qy 1194 ALSIGDKTHQFORSVSAKREALVGNPPIYRFKQNLQHQDSSVNVGTARWETAY 1253

Db 1236 alalallikldfsdsvpvrwnderyyggygstqatfavfagalagradvphkldnmd 1295

Qy 1254 ALLTSIMKIDINYNVPVIRKWLSEDRYGGGYSTQDTINAEGLFEYSILVKQLR-LSMD 1312

Db 1296 velhlparspvtvfillweegallrseetkqmegfeltak-gkgqgtlsvvtvyhakvkg 1354

Qy 1313 IDVSYKKGALANYKMT-DKNFLGRPVEILLNDDLLIVSTGFGSLATVHTVTVVHKTS 1371

Db 1355 ktctckldlrvtkpapatkppdaksimildicty-lg-dvdat-ms--ildismmt 1409

Qy 1372 EEVCS-FYLIKIDTDIEASHYRGYGNVDYKRVACASYKPSRESSSGSHAVMDISLPT 1430

Db 1410 gfipdtdnellsegvdrylskymekafsnkntliiykleishaeedclsfkvhqfnv 1469

Qy 1431 GTSANEDIKALVEGVQLFTDYQI-KD-GH---VIQIANSIPSSDFLCVRFRIFELEV 1485

Db 1470 gliqpskvkyvnylnleesctrfyhpkekddgmIsKlchmcrcaeeencf-mhqsgd-qv 1527

Qy 1486 GELSPATFTVYEHHRPKQCTMFY-S-TSNIKIQKVGCAACKCEADCGQMOEJLUTI 1543

Db 1528 elnerldkacepgvdyvytklttielsddfdeytmieqviksgsdevqagqerrfish 1587

Qy 1544 SAETRKQTAKEIAYAYKVSITSITVENFVKYKATLLDIYKTGEAVERKSEITFIKK 1603

Db 1588 vkcnalklqkqkylmwglss-dl-wgekptsyiigkdtwvehwp 1632

Qy 1604 VTCTNA-ELVKGRYLIMGKEALQIKNFESFRYIYPLDSLWTIEWP 1649

RESULT 9

ID C03\_EPTBU STANDARD; PRT; 1620 RA.

AC P98094;

DT 01-NOV-1995 (REL. 32, CREATED)

DT 01-NOV-1995 (REL. 32, LAST SEQUENCE UPDATE)

DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)

DE COMPLEMENT C3 (CONTAINS: C3A ANAPHYLATOXIN (FRAGMENT)).

GN C3.

OS EUPATRETUS BURGERI (INSHORE HAGFISH).

OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; PISCES;

OC AGNATHA (CYCLOSTOMATA).

RN [1]

RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.

RX MEDLINE; 92192016.

RA ISHIGURO H., KOBAYASHI K., SUZUKI M., TITANI K., TOMONAGA S., KUROSAWA Y.;

RL EMBO J. 11:829-837(1992).

CC -!- FUNCTION: C3 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE COMPLEMENT SYSTEM. AFTER ACTIVATION (C3B), IT CAN BIND COVALENTLY, VIA ITS REACTIVE THIOLESTER, TO CELL SURFACE CARBOHYDRATES OR IMMUNE AGGREGATES. CYCLOSTOMATES C3 APPEARS TO REPRESENT THE COMMON ANCESTOR OF MAMMALIAN C3 AND C4, SHOWING SIMILARITIES TO BOTH PROTEINS.

CC -!- SIMILARITY: TO C4, C5 AND ALPHA-2-MACROGLOBULIN.

CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.

DR EMBL; 211595; G62775; -.

DR EMBL; 211596; -; NOT ANNOTATED\_CDS.

DR PIR; S21045; S21045.

DR PROSITE; PS00477; ALPHA 2 MACROGLOBULIN.

KW COMPLEMENT PATHWAY; PLASMA; INFLAMMATORY RESPONSE; GLYCOPROTEIN; SIGNAL.

KW NON TER 1 1

FT CHAIN <1 1620 COMPLEMENT C3.

FT CHAIN <1 633 BETA CHAIN (BY SIMILARITY).

FT CHAIN 634 1336 ALPHA CHAIN (BY SIMILARITY).

FT CHAIN 1343 1620 GAMMA CHAIN (BY SIMILARITY).

FT PEPTIDE 634 714 C3A ANAPHYLATOXIN (BY SIMILARITY).





Qy 831 YSVRGQIQLGKTVYNYRTSGMFCVKMSAVEGICTSESPVIDHQGTSKCVQRKVEG 890  
 Db 871 mstrvpyviipmklglshiesvkaevknsgndgkdrllrvvaegvllvkketnvlhnp-v 929  
 Qy 891 SSSHLVFTVPLPLEJGJHNF--SLETFGKEILUWTLVWPEGVKRESYSQVTLDRG 948  
 Db 930 khgg-eqtshpvgprnqpsdadtllsvtagedevlveqalsgslvqpgvc 988  
 Qy 949 IYGTISRREPPYRIPIDIVPKTEKRLSVK-GLVGEILSNVLSQEGINILTHLPKGS 1007  
 Db 989 geqmiiymtlpviat-hyldntkwwedi-q--ldkrntalkviniqyqrlayrkedqsy 1044  
 Qy 1008 AEAEIMSV-VPVYVHYLETGNHNTIFHSDPLIEKOKLKKKREGMLSTMNRADYSY 1066  
 Db 1045 aawsvrqsatltayvvkvfamsstllisqenvlctavkwlilntqpdqglnfapvih 1104  
 Qy 1067 SVKRGGSASTLTAFALRVLCQVKNKYVEQONSICNSLLWLVNTYQLDNGSFKENSQYOP 1126  
 Db 1105 aemtgn--vrgsdadamtavfllamqeaasvceqsvslpagsmakavayl-ekrlp-h- 1159  
 Qy 1127 IKLAGTLPVEARENSLYTAFVIGIRKAFDICPL-VK-IDTALLIKADNELLTLETPAQS 1184  
 Db 1160 ---lt-npyavam--taya-l-anaglnketllkfaap-----q-lbh--wvpg-gy 1201  
 Qy 1185 FTIATISATSLGCKTHPQFISVSLAKREALVKNPPIYRFHKNLQHKUSSVPNTGT 1244  
 Db 1202 qytleatsvallalvkvfaeeagpivrnkqkvqvgvgstqetlmvfqavaevshv 1261  
 Qy 1245 ARMVETAYALLTSANIKDINTVAPVKMLSEEQRYGGGYSTQDTTNAIEGLEYSLLV 1304  
 Db 1262 kllkdfdmnlvagrastvwnnkqfhttrtdkvnslkdlitvkae-gnqeatslv 1320  
 Qy 1305 KQLR-LSMDIDVSYKHKAGLHNYKMTDN-FLGRPVEVL-LNDDLIVSTGSGLATVHV 1361  
 Db 1321 vlyyalpeeksdcesfdlvtltkmdktshd-akesfmltie-vlykns--erda-t 1375  
 Qy 1362 TTVVHTSTSEEV-C-SFYLIKIDQIDIEASHYRGVGNSDYKRIVACASYKPSRESSSGS 1419  
 Db 1376 m-sildlglitgftvtdldnglekgreryiekfmdkvlsergellilyldkvhshkldr 1434  
 Qy 1420 SHAVMDISLPTGISANEEDLKALVEGVQDLFTDYQI-K--D-GHVLQNSIPSSDFLC 1474  
 Db 1435 iefkhrvovglvqaavsvyeyn-qkrvkvfhpqreggtlarlclgdvctcaesc 1493  
 Qy 1475 VRFRIFELFVGLSPATVTYVEHRPDKQCTMFY-STSN1-KIQKVEGACACKVCEADC 1532  
 Db 1494 e-mktgepdrv-q-rldkacagldvvykatvdesklthtdtytkidvli-hpqtde 1548  
 Qy 1533 GQMOEELDTISAEKTRQACKPEIAYKVSITSIVENVEVVKATL-LDIYKTEAV 1591  
 Db 1549 gveqndrfmglaycrealglnqkqymimgksedlhvrvedkgllyqkyvlgeqtwey 1608  
 Qy 1592 AEKDEITETIKKVTNA-ELVKGQVYLMGK-EALQ-IKYNFSFRYVLDLSLWIEY 1648  
 Db 1609 peqectardvrevclgidefinqitfqc 1638  
 Qy 1649 PRDTTCSS--COAFLANLDEAFEDIFLNGC 1676

RESULT 11  
 ID C04 HUMAN STANDARD; PRT; 1741 AA.  
 AC P01028;  
 DT 21-JUL-1986 (REL. 01, CREATED)

DT 01-NOV-1990 (REL. 16, LAST SEQUENCE UPDATE)  
 DT 01-OCT-1996 (REL. 34, LAST ANNOTATION UPDATE)  
 DE COMPLEMENT C4 PRECURSOR (CONTAINS: C4A ANAPHYLATOXIN).  
 GN C4A AND C4B.  
 OS HOMO SAPIENS (HUMAN).  
 OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;  
 OC EUETHERIA; PRIMATES.  
 RN [1]  
 RP SEQUENCE OF 1-22 AND 1056-1225 FROM N.A.  
 RX MEDLINE; 85156269.  
 RA BELT K.T., YU C.Y., CARROLL M.C., PORTER R.R.;  
 RL IMMUNOGENETICS 21:173-180(1985).  
 RN [2]  
 RP SEQUENCE OF 20-1741 FROM N.A.  
 RC TISSUE=LIVER;  
 RX MEDLINE; 84156544.  
 RA BELT K.T., CARROLL M.C., PORTER R.R.;  
 RL CELL 36:907-914(1984).  
 RN [3]  
 RP SEQUENCE OF 680-756.  
 RX MEDLINE; 81264286.  
 RA MOON K.E., GORSKI J.P., HUGLI T.E.;  
 RL J. BIOL. CHEM. 256:8685-8692(1981).  
 RN [4]  
 RP SEQUENCE OF 957-1044.  
 RX MEDLINE; 82182029.  
 RA CAMPBELL R.D., GAGNON J., PORTER R.R.;  
 RL BIOCHEM. J. 199:359-370(1981).  
 RN [5]  
 RP SEQUENCE OF 990-1037.  
 RX MEDLINE; 82150875.  
 RA HARRISON R.A., THOMAS M.L., TACK B.F.;  
 RL PROC. NATL. ACAD. SCI. U.S.A. 78:7388-7392(1981).  
 RN [6]  
 RP STRUCTURAL BASIS OF POLYMORPHISM.  
 RX MEDLINE; 87080272.  
 RA YU C.Y., BELT K.T., GILES C.M., CAMPBELL R.D., PORTER R.R.;  
 RL EMBO J. 5:2873-2881(1986).  
 RN [7]  
 RP VARIANT C4A6 ALLOTYPE.  
 RX MEDLINE; 92242905.  
 RA ANDERSON M.J., MILNER C.M., COTTON G.H., CAMPBELL R.D.;  
 RL J. IMMUNOL. 148:2795-2802(1992).  
 CC -!- FUNCTION: C4 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE  
 CLASSICAL PATHWAY OF THE COMPLEMENT SYSTEM. IT IS PROCESSED BY  
 ACTIVATED C1 WHICH REMOVE FROM THE ALPHA CHAIN THE C4A  
 ANAPHYLATOXIN.  
 CC -!- SUBUNIT: THIS PROTEIN IS SYNTHESIZED AS A SINGLE-CHAIN PRECURSOR  
 AND, PRIOR TO SECRETION, IS ENZYMATICALLY CLEAVED TO FORM A TRIMER  
 OF NONIDENTICAL CHAINS (ALPHA, BETA, AND GAMMA).  
 CC -!- POLYMORPHISM: HUMAN COMPLEMENT COMPONENT C4 IS POLYMORPHIC WITH AT  
 LEAST TWO LOCI, C4A & C4B. 13 ALLELES OF C4A & 22 ALLELES OF C4B  
 HAVE BEEN DETECTED. THE ALLELE SHOWN HERE IS C4A4.  
 CC -!- C4A ALLOTYPES REACT MORE RAPIDLY WITH THE AMINO GROUP OF PEPTIDE  
 ANTIGENS WHILE C4B ALLOTYPES REACT MORE RAPIDLY WITH THE HYDROXYL  
 GROUP OF CARBOHYDRATE ANTIGENS.  
 CC -!- POLYMORPHISM: THE C4A ALLELES CARRY THE BLOOD GROUP RODGERS WHILE  
 THE C4B ALLELES CARRY THE BLOOD GROUP CHIDO.  
 CC -!- DISEASE: THE C4A6 ALLOTYPE IS TOTALLY DEFICIENT IN HEMOLYTIC  
 ACTIVITY.  
 CC -!- FUNCTION: DERIVED FROM PROTEOLYTIC DEGRADATION OF COMPLEMENT C4,  
 C4A ANAPHYLATOXIN IS A MEDIATOR OF LOCAL INFLAMMATORY PROCESS. IT  
 INDUCES THE CONTRACTION OF SMOOTH MUSCLE, INCREASES VASCULAR  
 PERMEABILITY AND CAUSES HISTAMINE RELEASE FROM MAST CELLS AND





Qy 1231 NIQHD-S-S-VP-N-TGT---AR-M-VETTAIALTSL-NIKDINTVNPVNIKWSEQR 1279

Db 1295 fggfstqdtvialdaelayiahtteerglnvltsetgrngfkshalqimnrqirgl 1354

Qy 1280 YGGFYSTODTINAIEGLTEYSLL--VKQIR-LSMDID-VS---YK-HKCALFN--YK-M 1328

Db 1355 eelqfslgskinvkvngskgtikvlrtnvldmktctqdlqievtkghvetymean 1414

Qy 1329 TDK-NF-LGRPVLL--ND-D-LIVSTGFGS-GL--AT---VHV-TTV-VHKTSTSEEV 1374

Db 1415 edveydelpakddpqlpvtplqlfegrrnrrreapkvveeqesrvhvtcvirngk 1474

Qy 1375 CSF-Y--L--KIDTQD-IEA-SHRGCV-GNSDYKR-----IVA---C-ASYKPS-REESS 1416

Db 1475 vqlsmaiadvtlllegfhalradlektsldryvshfetegphvllfydsvpter-ecv 1533

Qy 1417 SG-SSHAVNDISLPTGISANEEDLKALVEGVQDLFTDYQIKDGHVILQINISIPSSDFLCV 1475

Db 1534 gfeavqevpvlvpasatllydynperrcsfygapkskerlllatlcsaevcqaegtcp 1593

Qy 1476 RERIFELFEVGLSPATFTVVEYHRPDKQCTMFYST-SNIKI-QKVCEGAACKVCEADCG 1533

Db 1594 tqralergldedgymkfacyprvevfgvkvredraafrlfetkitqvlhftkd 1653

Qy 1534 QMQEELDITI-SAET-RKQTACK-PEIAYAYKVSITSITVENFVKYKATLDDIYKTGEA 1590

Db 1654 vkaanqmrnfivraec-frlrlepkeylimlqdg--atydlghpqyldlenswiemp 1710

Qy 1591 VAEKDSEI-TPIKKVCTTNAELVKGROYLIMCKEALQIKYNSFRYIYPLDSLTWIEWP 1649

Db 1711 eerlcrstrqaacqndflqeygtgc 1739

Qy 1650 RDTTC-SSCQ-AFLANLDEAFIDFNGC 1676

RESULT 12

ID C04 MOUSE STANDARD; PRT; 1738 AA.

AC P01029;

DT 21-JUL-1986 (REL. 01, CREATED)

DT 01-AUG-1988 (REL. 08, LAST SEQUENCE UPDATE)

DT 01-OCT-1996 (REL. 34, LAST ANNOTATION UPDATE)

DE COMPLEMENT C4 PRECURSOR (CONTAINS: C4A ANAPHYLATOXIN).

GN C4.

OS MUS MUSCULUS (MOUSE).

OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;

OC EUTHERIA; RODENTIA.

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE; 85298264.

RA SEPICH D.S., NOONAN D.J., OGATA R.T.;

RL PROC. NATL. ACAD. SCI. U.S.A. 82:5895-5899(1985).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=B12.WR;

RX MEDLINE; 87309760.

RA ROSA P.A., SEPICH D.S., ROBINS D.M., OGATA R.T.;

RL J. IMMUNOL. 139:1568-1577(1987).

RN [3]

RP SEQUENCE FROM N.A.

RC STRAIN=B12.WR; TISSUE=LIVER;

RX MEDLINE; 89380278.

RA OGATA R.T., ROSA P.A., ZEPF N.E.;

RL J. BIOL. CHEM. 264:16565-16572(1989).

RN [4]

RP SEQUENCE FROM N.A.

RC STRAIN=FW; TISSUE=LIVER;

RX MEDLINE; 85289294.

RA NONAKA M., NAKAYAMA K., YEUL Y.D., TAKAHASHI M.;

RL J. BIOL. CHEM. 260:10936-10943(1985).

RN [5]

RP SEQUENCE OF 651-810 AND 924-1083 FROM N.A.

RX MEDLINE; 85038607.

RA NONAKA M., TAKAHASHI M., NATSUUME-SAKAI S., NONAKA M., TANAKA S.,

RA SHIMIZU A., HONJO T.;

RL PROC. NATL. ACAD. SCI. U.S.A. 81:6822-6826(1984).

RN [6]

RP SEQUENCE OF 1105-1449 FROM N.A.

RX MEDLINE; 85166208.

RA LEVI-STRAUSS M., TOSI M., STEINMETZ M., KLEIN J., MEO T.;

RL PROC. NATL. ACAD. SCI. U.S.A. 82:1746-1750(1985).

RN [7]

RP SEQUENCE OF 1257-1376 FROM N.A.

RX MEDLINE; 85038859.

RA TOSI M., LEVI-STRAUSS M., DUFONCHEL C., MEO T.;

RL PHILLOS. TRANS. R. SOC. LOND., B, BIOL. SCI. 306:389-394(1984).

RN [8]

RP SEQUENCE OF 1360-1511 FROM N.A.

RX MEDLINE; 83273751.

RA OGATA R.T., SHREFFLER D.C., SEPICH D.S., LILLY S.P.;

RL PROC. NATL. ACAD. SCI. U.S.A. 80:5061-5065(1983).

RN [9]

RP SEQUENCE OF 1-128 FROM N.A.

RC STRAIN=FW; TISSUE=LIVER;

RX MEDLINE; 86031969.

RA NONAKA M., NAKAYAMA K., YEUL Y.D., SHIMIZU A., TAKAHASHI M.;

RL IMMUNOL. REV. 87:81-99(1985).

CC -!- FUNCTION: C4 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE

CC CLASSICAL PATHWAY OF THE COMPLEMENT SYSTEM. IT IS PROCESSED BY

CC ACTIVATED C1 WHICH REMOVE FROM THE ALPHA CHAIN THE C4A

CC ANAPHYLATOXIN.

CC -!- SUBUNIT: THIS PROTEIN IS SYNTHESIZED AS A SINGLE-CHAIN PRECURSOR

CC AND, PRIOR TO SECRETION, IS ENZYMATICALLY CLEAVED TO FORM A TRIMER

CC OF NONIDENTICAL CHAINS (ALPHA, BETA, AND GAMMA).

CC -!- C4 IS A MAJOR HISTOCOMPATIBILITY COMPLEX CLASS-III PROTEIN.

CC -!- SIMILARITY: TO C3, C5 AND ALPHA-2-MACROGLOBULIN.

CC -!- SIMILARITY: CONTAINS ONE ANAPHYLATOXIN-LIKE DOMAIN.

DR EMBL; K00019; E19386; -.

DR EMBL; M11729; G387420; -.

DR EMBL; M12968; G199270; -.

DR EMBL; M12970; G199262; -.

DR EMBL; M12972; G199264; -.

DR EMBL; M1789; G387438; -.

DR EMBL; K02798; G199282; -.

DR EMBL; M17440; G387440; -.

DR PIR; A01264; A01264.

DR PIR; A21692; A21692.

DR PIR; A2039; A2039.

DR PIR; A24558; A24558.

DR PIR; A29059; A29059.

DR HSP; P01032; I65A.

DR PROSITE; PS00477; ALPHA 2 MACROGLOBULIN.

KW COMPLEMENT PATHWAY; PLASMA; GLYCOPROTEIN; MHC III; SIGNAL;

KW INFLAMMATORY RESPONSE.

FT SIGNAL 1 19

FT CHAIN 20 673 BETA CHAIN.

FT CHAIN 678 1443 ALPHA CHAIN.

FT CHAIN 1448 1738 GAMMA CHAIN.

FT PEPTIDE 678 753 C4A ANAPHYLATOXIN.

FT DOMAIN 700 734 ANAPHYLATOXIN-LIKE.  
 FT DISULFID 700 726 BY SIMILARITY.  
 FT DISULFID 701 733 BY SIMILARITY.  
 FT DISULFID 714 734 BY SIMILARITY.  
 FT THIOLIST 1006 1009 BY SIMILARITY.  
 FT CARBOHYD 224 224 POTENTIAL.  
 FT CARBOHYD 743 743 POTENTIAL.  
 FT CARBOHYD 1387 1387  
 FT CONFLICT 132 132 F -> Y (IN REF. 4).  
 FT CONFLICT 327 327 G -> E (IN REF. 4).  
 FT CONFLICT 570 570 Q -> E (IN REF. 4).  
 FT CONFLICT 720 720 R -> G (IN REF. 5).  
 FT CONFLICT 739 740 DL -> AI (IN REF. 5).  
 FT CONFLICT 838 838 P -> R (IN REF. 4).  
 FT CONFLICT 993 993 P -> L (IN REF. 5).  
 FT CONFLICT 1043 1043 D -> E (IN REF. 5).  
 FT CONFLICT 1119 1119 V -> A (IN REF. 6).  
 FT CONFLICT 1190 1190 A -> T (IN REF. 6).  
 FT CONFLICT 1324 1324 K -> N (IN REF. 4).  
 FT CONFLICT 1401 1401 G -> S (IN REF. 8).  
 FT CONFLICT 1442 1442 R -> K (IN REF. 4).  
 FT CONFLICT 1453 1453 A -> V (IN REF. 4).  
 SQ SEQUENCE 1738 AA; 192870 MM; A6CCIFCB CRC32;  
 Query Match 13.3%; Score 1605; DB 2; Length 1738;  
 Best Local Similarity 27.6%; Pred. No. 0.00e+00;  
 Matches 449; Conservative 407; Mismatches 628; Indels 143; Gaps 111;  
 Db 137 gbiwtdpypgqrvrvyrfaldqmpetdfilitvsnghlrv-lkkel-fsts 194  
 Qy 124 GFLEIHTDKPVYTDQSVKRVYSLNDLKPAGRETVLTFIDPGESEVDNVEIDHIGII 183  
 Db 195 ifqdaftidiseptgwksarfdglsnsthfevkvipnfevktpwkpilmvp 254  
 Qy 184 SFPD-FKIPSNPRYGMWTKAKYKEDFSTGTAYFVKEVYLPHFVSIEP-E-YNFIGY 240  
 Db 255 snsdeldiqdaryigkpv-qgvaytrfalmdc--qgkrtflrqltqaklvegrthis 311  
 Qy 241 KMFKEITIKARFYKVKVTEADVYITFGIEDIKDDQKEMQMTAMQNTMLINGIAQVT 300  
 Db 312 lskdqqaalkdnigrvdeglrlyaataviespggeneaealtswrfvsaafeldslr 371  
 Qy 301 FDESETAVKELS--YYSLEDJANKLYLTAIVTVIESTGCFSEEAETPGIKYVLSPYKLINVA 358  
 Db 372 ttrhlvpqghflqalvqemsgseasnrvpkvsa-tlvsgdsqvldiqstngiqq-v- 428  
 Qy 359 TPLFLKGPPIPKVQVKDSLQVLGCVPLVNAQTIDVNETSDLPSSKSVTRVDDGVA 418  
 Db 429 eisfpipptvtelrlvsagel-ygal--arltvga-p-psrtgfgflsie-pldpresv 482  
 Qy 419 SEVLNLPSCVTVLEFNKVDAPDLPENQARECYRAIAYSSLSQSYLYIDWDHKKALLV 478  
 Db 483 gdtfiinlqpvqipaptfshyvvymilergimang-rep-rktv-tsavvlvdhqlapsf 539  
 Qy 479 GEHLNIIVTPKSPYDKITHYNYLILSKGKIIFHGTREKFSQASYSINIPVQNWPPSS 538  
 Db 540 yfvayfyhag-h-p--vanslliniqerdcgkqlkvd-qakeyrnadmmlkriqtcdsk 594  
 Qy 539 RLLVYIVTGEQTAELVSDSVMLNIEEK-CGNQLQVH! SPDADAYSPGQVTSVNNATGMD 597  
 Db 595 alvalqavdtaiyavgrgshkpldmkskvfevinyngvpggddalqvfdaglafsd 654  
 Qy 598 SKWALAAVDSAVYGVQRGAKRPLE--RVQFQLEKSDIGCCAGGLNNANVFHLAGLTLT 655

Db 655 gdrltqtre-dlscppekksrqtrnmvfkavseklqgyespdakrcqqdmtkklpmkrt 713  
 Qy 656 NANADDSOENDEPC-KEIL-RPRTL--QKKIEEIAAKYKHSVVKKCCYDGCAC-VNNDET 710  
 Db 714 ceqraarvpqqa-crepfiscckfaedlrrnqtrsqahlarhnmhmlqeedlideddliv 772  
 Qy 711 CEQRAARISIGPRCIKAFTECCVVASQIRANISHKDMQLGRL-H-K--TLIPVSKPEI 765  
 Db 773 rtsfpenwlrvepvdesklitwlpdsmttwihgvskskglcvakptrvrvfrkfh 832  
 Qy 766 RSYFESLWELVEHLVPRRKQJQAFALPDSLTTWETIGTIGISNT-GICVADTVKAKVFOVF 824  
 Db 833 lhlrlpisirrfefelrpvlynylnndvavshvrtvpeglclagggmmaqqvtvpagasa 892  
 Qy 825 LEWNIPIYSVRGEQIQLKCTVYNYRTSQMFCVKMSAVEGICTSESPVIDHQTKSKCV 884  
 Db 893 rp-vafsvvptaaanv-plkvvargv-fdl----g-davekilqiekaghaihreeiv-yn 943  
 Qy 885 RQKVEGSSHLVTFVTLPLEIGLHNINFSLETWFKGKELVKTILRVVPEG-VKRESYSGVT 943  
 Db 944 ldp--l-nnlgrtllpigsdgnivpdqdfsslvrvrtasepletmgsegalspggvaell 1000  
 Qy 944 LDPGICYTISRKEFFYPIDILVPKTEKRLISVKGLLVGEIL-S-AVLSEGINILT 1001  
 Db 1001 lrpqgaetmiylapltasnyldrtewsklspet-kd-havdliqk-gymriqqfrk 1057  
 Qy 1002 HLPKSAEALMSVVPVYVFHYLETGNHNIHFSDPLTEKOKLKKLKEGMLSTMSYRN 1061  
 Db 1058 ndqgfawlhrrdsrwtatfvikilsaqeqvgnspesklqetaswlaq-qlqdgshdp 1116  
 Qy 1062 ADYSYSVWKGSASTWLTAFALRVLGQWKYVEQONQNSICNSILLWLVENYQLDNGSFEN 1121  
 Db 1117 cpvlhramgggl-vgs--detvaltafvialhhlhldvfdqddakqlknrveasaitkansf 1174  
 Qy 1122 SOYQPKLOGTLPVEARENSLYLTAFTVIGIRKA---F--DIC-PL-VKIDTALIKADNF 1174  
 Db 1175 lqkaseallgahaaaitayaltit-kaseelrlnrvahnslmamaetgeh-lywglvlgs 1232  
 Qy 1175 LIENLPAQ-STFTLISAYALSGLDTHPQFRSIV-SALKREALVKGNPPIYR---F-W 1228  
 Db 1233 qdkvvlrptapreptepvpqapalwiettayall-hllireqkgkmadkaaswlthqgsf 1291  
 Qy 1229 KDN--LQHKDSSVP-NTGT-AR-M-VETTAYALLTSJNLKD-I-NYVNPVIRWLSEEQRY 1280  
 Db 1292 hqafretqdtvrtldalaywiashtteekalkvtlsamgrnglktghlhlhnhqvkgle 1351  
 Qy 1281 GGGFYSTQDTINAEGLTEY---SLIV--KQLRLSMD-ID-VSYKHKA-LHNY--K-MT 1329  
 Db 1352 eelfelgetisvkvneqskgtklirtnvnlmdknttcqdlqievkvgtgaveyawardane 1411  
 Qy 1330 DK-NF-LGRPEVIL--ND-D-L-IVST-G-FG----S--GLAT-VHVTTVHKT-STSE 1372  
 Db 1412 dyedydpmaadpsvplqpvtpqlfegtrrrrrreapkaeqesrvqyvtciwrngk 1471  
 Qy 1373 EVCSFY-LKI-DTODI--EA-SHYRGY-QN-SDYKR-I---VA----C-ASYKES-RESS 1416  
 Db 1472 lqlagmaaditllqghalradlekltlsldryvshfetdphvlllyfdfspttr-ecv 1530  
 Qy 1417 SG-SSSHVMDISLPTGISANEEDLALVEGVQDLFTDYQIKDCHVILQINSIPSSDFLCV 1475  
 Db 1531 gfsgaeqsvrvlqvpsavlydyyspdhtcsvfyaaptkaqlilatlcsgdvccqcaegkcp 1590  
 Qy 1476 RFRIFELFEVGLSPATFTVVEYHRPDQCTMFYST-SNIKI-QKVCEGAACKCVCEADCG 1533



Db	1591	rllrelervedkdyrmfacyprvveygtvkvlredgraarlfeskitqyvlhfrkd	1650
		Q:   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :	
Qy	1534	QMOEELDTISA-ET-RKQTACK-PEIAYAKVYSITSITVENFVKYATLLDIYK-TGE	1589
		Q:   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :	
Db	1651	tnasigtgrnflrasc-rllrlepkeylvimgndgetednkgdpqyl--ldsnrtwiemp	1707
		Q:   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :	
Qy	1590	AVAEKDSSEITFKKVTCTNAELVKGRQYLVLMGKEALIKYNF5FRYIYPLDLSLTWYMP	1649
		Q:   :   :   :   :   :   :   :   :   :   :   :   :   :   :   :	
Db	1708	seqmcks	1714
		:	
Qy	1650	RDTTCSS	1656

RESULT	13
ID	C03 RABIT
AC	P12247;
DT	01-OCT-1989 (REL. 12, CREATED)
DT	01-OCT-1989 (REL. 12, LAST SEQUENCE UPDATE)
DT	01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)
DE	COMPLEMENT C3 ALPHA CHAIN (FRAGMENT).
GN	C3.
OS	ORYZOLAGUS CONICULUS (RABBIT).
OC	EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
OC	EUTHERIA; LAGOMORPHA.
RN	[1]
RP	SEQUENCE FROM N.A.
RX	MEDLINE: 87006907.
RA	KUSANO M., CHOI N.H., TOMITA M., YANAMOTO K., MIGITA S., SEKIYA T.,
RL	NISHIMURA S.;
RR	IMMUNOL. INVEST. 15:365-378(1986).
CC	- - FUNCTION: C3 PLAYS A CENTRAL ROLE IN THE ACTIVATION OF THE COMPLEMENT SYSTEM. ITS PROCESSING BY C3 CONVERTASE IS THE CENTRAL REACTION IN BOTH CLASSICAL AND ALTERNATIVE COMPLEMENT PATHWAYS. AFTER ACTIVATION C3B CAN BIND COVALENTLY, VIA ITS REACTIVE THIOLESTER, TO CELL SURFACE CARBOHYDRATES OR IMMUNE AGGREGATES.
CC	- - SUBUNIT: C3 PRECURSOR IS FIRST PROCESSED BY THE REMOVAL OF 4 ARG RESIDUES, FORMING TWO CHAINS, BETA & ALPHA, LINKED BY A DISULFIDE BOND. C3 CONVERTASE ACTIVATES C3 BY CLEAVING THE ALPHA CHAIN,
CC	RELEASING C3A ANAPHYLATOXIN & GENERATING C3B (BETA CHAIN + ALPHA' CHAIN).
CC	- - SIMILARITY: TO C4, C5 AND ALPHA-2-MACROGLOBULIN.

DR	PROSITE; PS00477; ALPHA 2 MACROGLOBULIN.
DR	PROSITE; PS00477; ALPHA 2 MACROGLOBULIN.
KW	COMPLEMENT PATHWAY; COMPLEMENT ALTERNATE PATHWAY; PLASMA;
KW	INFLAMMATORY RESPONSE; GLYCOPROTEIN.
FT	NON TER 1 1
FT	CHAIN <1 726 ALPHA CHAIN.
FT	THIOLEST 73 76
FT	CARBOHYD 2 2 POTENTIAL.
FT	CARBOHYD 233 233 POTENTIAL.
FT	CARBOHYD 680 680 POTENTIAL.
SQ	SEQUENCE 726 AA; 81844 MW; 27A509CF CRC32;

Query Match 6.6%; Score 796; DB 2; Length 726;  
Best Local Similarity 26.2%; Pred. No. 5.45e-148;  
Matches 197; Conservative 197; Mismatches 305; Indels 52; Gaps 37

Db	9	tldpenlqggvy-qkeipsadisqdyptesetkil-lqgtvpvqmteidaiderlhl	66
		:   : :    :    :    :    :    :    :    :    :    :    :	
Qy	943	TLDPRGI-YGTISRKEFFYPRIIDLVPKTEI-KRLISVKGLLVGSEILSAVLSEQENIL	1000
Db	67	ivrgscgcgmiamthvtviavhyldhteqdxf-s--lekqealelikkgytqqafsk	123
		:      :      :      :      :      :      :      :      :	

QY	1001	THLPKGSAAELMSVVPVFVTFYHLETGNHNIFHSODPLTEKORUKKKLKEGMLISMTSR	1068
Db	124	qpnosyaaflnrapetwltayvkvfslawnlliaidsqvlgcgvkwlmeakqpdgvyfce	183
QY	1061	NADYSYSVMKGSGASTWLTAFALRVLGQWKVYEQNQSICNSLWLVENYQLDNGSFKE	1120
Db	184	dapvlhqem:gggr-nsekeralkafvylalqualgeareiceeqvnslaasin kardflan	242
QY	1121	NSOYQPIKLQGLTPVEARENSLVLTAFVVICIRKAFDICP-LVK-IDTALKADNLEEN	1178
Db	243	ymlnlprpyvaiaawaqqdk----lrg--aflnk-flskakek-nr-weeppqr-?----	289
QY	1179	TIPAQSTFFILISAIALSLGBDKTHPQRSIVSALKREALVKGNPPYFRFMDNLQHWOSS	1238
Db	290	l-yn-----veasyallailldrfdsvpvrwlneqryyggvgstqatmgfqala	343
QY	1239	VPTCTARWETTAYALLTSANIKOINYNPVIMKLSSEORYGGGYSTQDTINAIEGLT	1298
Db	344	qvctdvdphkdlnmvwsiqlpersspvkhrivwdsasllrseetkenggfeltacq-gkgqc	402
QY	1299	EYSLAVKQLA-LSMIDIVSYKHKGALHNYKWT-DKNFLGRPEVELINDDLIVSTGFSGSL	1356
Db	403	qlgwrttyfakvkvctckfdrlvniktapevkpqdakemilghctrj-lq-ded	460
QY	1357	ATVHVTTVMKISTEEVCVS-FYLKDITQDIASHRYCGNSDYKRIVACASVKPSREES	1415
Db	461	at-ma---ildismgtfvdpdtddlnlletgdvtryiskyeInkafsenkntliijldkshs	517
QY	1416	SSGSHAVMNDISLPTGISANEEDIKALVEGDQGLTDYQT-KD-GH---VIIJLANSTPSS	1470
Db	518	reclafakhvqfnvglipgavkyvsymleetctqfyhekedmksklchkmcrca	577
QY	1471	DFLCYRFRIFELLEVEFGLESPATFTVYEHPRDKQCTMEY-S-TSNIKIQKVCEGAACKCV	1528
Db	578	eencf-mq-q-ldekittndrldkacepldgvyvtktlvqveradddfeylmvventiksg	635
QY	1529	EADCQQOELEDLITSNETRKQTACKAEIAYAKVISITSITWENVFYKYKATLLIDYKTG	1588
Db	636	sdevqagqpafishikrcdaikldkgkhylmwglsdpvqeKptnsylvigkd--twvef	693
QY	1589	EAVAEXOSEITFIKKVTCYNA-ELVKGROYLINGKEALQIKNFYRIYPLDSLTMWE	1647
Db	694	wpekeecdeeqnhcedlga faesmvmvcg	724
QY	1648	WPRDPTC-SS-QCAF-LANDIDEADEFINGLC	1676

RESULT	14	STANDARD;	PRT;	1477 AA.
ID	AL13 RAT			
AC	P14046;			
DT	01-APR-1990 (REL. 14, CREATED)			
DT	01-APR-1990 (REL. 14, LAST SEQUENCE UPDATE)			
DT	01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)			
DE	ALPHA-1-INHIBITOR III PRECURSOR.			
OS	RATTUS NORVEGICUS (RAT).			
OC	EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; MAMMALIA;			
OC	EUTHERIA; RODENTIA.			
RN	[1]			
RP	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.			
RC	TISSUE=LIVER;			
RX	MEDLINE; 88153707.			
RA	BRACIAK T.A., NORTHEMANN W., HUDSON G.O., SHIELDS B.R., GEHRING M.R.,			
RA	FEY G.H.;			
RL	J. BIOL. CHEM. 263:3999-4012(1988).			

Db	243	deini--vtacatytykvpv-ghvkief-chnpftfsetckegckeedsrldmngcstq	298
Qy	241	KNFNKFETIKARYENKVVTEADVITFGREDIKDOKEMQAMQMTWL--ING-IAQ	298
Db	299	evnitefqlke-ny--lk-mhqaf-hvnatvtteegtgefsagsgriever-trnkf-lfl	351
Qy	299	VTFDETAVKELSYSLIEDLNKKYLIYAVTVIE-STGG-FSEAEIPIGKIVLSPYKLN	356
Db	352	kadsnf-rhgifpfvkvrldlkgdpnqvlikardagyttnattddqhlakfaidtn	410
Qy	357	VATPFLFKGPIPIKVOVKDSLQJUGCVPVILNAQITDVNQETSDLDPSKSVTFVD-D	415
Db	411	giedyslknvykhkeescihascetaerhaeah-htay-av-ys-lsksyliid-tea-g	464
Qy	416	GVASFVNLDPSPGVTFLENVKTD-APDLPENOAEGVRAIYASLSOSYLIIDWTDHMK	474
Db	465	vlpcnqhtvqhafiikqvgvlgvllqvlfvlylvmaggsilqtgnthqvgpessvqgnf	524
Qy	475	ALLVGE-H-LNIVTPKSPYIDKITH--YNYLILSKGKLIHFG--TRE-KFSDASYQ--	524
Db	525	aleipwefsvvpmakmiylilp-d-geviadsvkfve-kcltn-kvhlfsfpqalp	579
Qy	525	SINIPVQNWPSSRLIVYITVGTQAEIIVSDVSLNIECKGNLQVHLIS--PDADAYS	583
Db	580	asqet-hmrvtasppqlcqlrvadvql-lq---kp-e-a-e-lsps-liydlpg-mqds	627
Qy	584	PGQTVSINMATGMDSWAALAAVDSAVVQVGQAKRPLERVQFLEKSDLGCGAGGGLANA	643
Db	628	n-f-las-s--ndpf-ed-edylmvgpiarek-dvyvrvretglmafnlkiklpty--	677
Qy	644	NVFLAGLITELINNAADDSQENDECKEILRPRTLQKLTKEELAAK-YKUSVVKCCYDG	702
Db	678	-cntdydmvplavpavaldestdgmysesipvva-vkepqpgepprkdpdp-kd--pvie	732
Qy	703	ACVNNDETCQRAARIISLGRPCIKAFTECCOVASQALRANISHKMQGLRILMKTLIPVSK	762
Db	733	t-lrnyfpetwldlvtvmsagvtelemtpdttewkagalcslndtqgliesvaeq	791
Qy	763	PEIRSYFESWLWE-VHLVPR-RKQJQFALPDSLTWTEIOGIGISN-TGICVADVTKAVK	819
Db	792	fpqpfwelampysvirgeaftlkatvlnylptslpmavllaeapdf-ta-vpennqd--	847
Qy	820	FRDVELEMNPIVSVRGEOIQLKGTVVNYRTSQMFCVKMSAVEGICTSESPVIDHQGTK	879
Db	848	sy-clangrhteswlvtpkqlg-nvnf-avs-a-eaqcpqpcgsevatvpctgrkdtv	902
Qy	880	SSKCVRKQVEGSSHLVTFVPLPLEIGHNINFSLETWFGKEILVKTLRVVPEGVKRESY	939
Db	903	xkvliivepegiikehtfssllcasdaelsetlslpbtvvdksarahfvmgdlisai	962
Qy	940	SvV-TLDPGCI-----YGTISRRKEFPVRIIDLIV-PKTEIKRILSVKGLLVEITLSAVL	992
Db	963	-kntqn-liqmpygcgeqnmvlfnapiyvlkyln-etq-ql--teklkaka-lgy-lrag	1014
Qy	993	SOEGINILTHLPKGSAAEIMSVVPVVFVHYELTGNHWNIFHSODPLEKQKLKKIKLEG	1052
Db	1015	yqrelnykhkdgysaafqdhngggngntwtatfvlksfaqarafideshitdaftwls	1074
Qy	1053	MUSIMSTRNAD--YS-YSVWKQ-GSASTWITAFALRVJGVQNKVYEQNQNSIONSLLMLV	1108
Db	1075	kq-qkdsqcfressagellnmakggvddetilaav-yit-mallesslp-dtdpvvskalec	1130
Qy	1109	ENYQJLNDGSPKSNQYQIKQACTLIPVEARENSLYLTAFTVIGIRKAFDPCPLWKIDTAL	1168

411	qisdyslnikyvhkeesc:ihsec:taerheaah-htay-av-ys-lksyiyid-tea-q	464
416	GVASFVILNPGSGVTLEFNWKT-DAPDLPEENOAREGYATAYSSLSQYIYIDTMDNHK	474
465	vlpcnqihqtqahfikiqvglvglqvlfvhyhlvmsagailqcmhthqvpesgvqgnf	524
475	ALLUGE-H-LNIIVTPKSPYIDKITH--YNYLISKGKGIHF-G--TRE-KFSDASYY--	524
525	aleipfsvmsvpvkmlyitlp-d---geviadsvkfve-kcltn-kvhlfspsqelp	579
525	SINIPVQNWPSSRLVYIVTGTQAEIUSVSWLNIIEKQNLQVHLS--PDADAYS	583
580	aegct-hmrvtasqqlcqlravdqsvl-lq---kp-e-a-e-lsps-liydipg-qmds	627
584	PQGVTSINMATGMSWALNADVSNAVYCVORGAKXPLERVFOLEKSDLCGGAGGLNNA	643
628	n-f-las-s--ndpf-ed-edyclmlygpiarek-dwyvvtretqlmafnlkikpty--	677
644	NFHLAGLITFLNNAADSGENDECKEILRPRTLQXKITEEIAAK-YKUSVVRKCCVDG	702
678	-cntdydmvlpavaldsdtgmwyeslpvva-vksploqeporkdppp-kd--pvie	732
703	ACVNNDTEQZRAARIISLGRPCIKAFTECCCVASQALRANISHKQWLGRILHMKTLLPYSK	762
733	t-irnyfpetwldvtvmsagvtelemtpdtitewkagalcndntgiglesvaeq	791
763	PEIRSYFESPLWE-VHLVPR-RKQALFALPDSLTTWIEIGIGISN-TGICVADVTYKAKV	819
792	fqpfvwlampysvirgeaftkatvlnytpslpmauvllaeapdf-ta-vpvennqd--	847
820	KFOVPELWNPTYSVVRGEQIQLKGTVYNYRTSQMOFCVKMSAVEGICTSESPVIDHQTK	879
848	sy-clqanghrteswlvtpkqlg-nvnf-avs-a-eaqspgpcgsevatvpctgrkdtv	902
880	SSKCVRKQVEGSSHLVTFVTLPLEIGHINFSLETFWFKELIWKTLRVVPEGVKRESY	939
903	kvkvlivepegiikentfssllcasdaelsetlslpbtvvdksarahsfvmgdilseai	962
940	SzV-TLDPRCI-----YGTISRRKEFPVRPJDIIV-PKTEIKRILSVKGLLVEILSAVL	992
963	-kntqn-llqmpygqcgemvlfapniyvklyn-etq-ql--tekikaka-lgy-lrag	1014
993	SEGINILITLTPKSAEAEIEMSVVPVVFVHYLETGNHWNIFHSOPLEIKQKLKKIKEG	1052
1015	yqrelnykhkdgysaafgdhngggngntwtatvfksfaqarafideshitdaftwls	1074
1053	MUSIMSTRNAD--YS-YSVWK-GSASTWITAFALRVJGQVKNYVEQWQNSIONSLLMLV	1108
1075	kq-qkdsqcfresagellnmamkggyvdeitlaa-vit-mallesslp-dt-dpvvekalcc	1130
1109	ENYQJLNDGSPKENSQYQIKQICTLPVARENSLYITAFVTGIRKAFDTCPLWIKIDTAL	1168

Db	1131	lesswienieggnfgyvfkalmayafalgagnckr--neilksldkea--ikednsiwher	1188
Qy	1169	IRADNFI LLENTLPAQSTFTLAI SAVALSISGDKTHQFQRSIVSALKREALKGNPNYYRWF	1228
Db	1189	pqkptkseeqlytpqasaeavemsayvlarltagpaspesdalemgitkwltkqmsy	1248
Qy	1229	KDNLQHKDSSV--PNGTGTARWETAYAA--L--LTSL---NLKDINYNPVVKMLSEQR--Y	1280
Db	1249	gg--festqdtvwaldalekyga--atfsksqtpstvsgsgsfqskfqvdkenlllqqv	1306
Qy	1281	GGGFTSTQDTINAI EGLTEYSLVLKQLRLSMDIDVSYKWKAL--HNYKMTDKN--FLGRPV	1338
Db	1307	slpypgnytvsvs--gegcvyag--tllcynvplekqgqapalqvtpvlctnmpkg--qns	1363
Qy	1339	EV--LUNDDLVITGFGSLGATVHTVVHKTSTSTSEEVCSFYLIKIDTQDIEASHYRGUNS	1397
Db	1364	--fajele--isyngst----pas-nm-viadvkmlsgfipktpvklk--erlighv--srtevt	1415
Qy	1398	DYKRIVACASYKPSREESSGSHVMDISLPTGTISANEEDJALVEGDQLFTDYOIKD	1457
Db	1416	nvvllyldqv--tnqtlsfaffiqddipvklqpaivkydyvtd	1459
Qy	1458	GWIIQLNISTPSDGLCYRRIRIFELFEVGLSPAFFTYVYTHRPD	1502

RESULT	15	STANDARD;	PRT;	1476 AA.
ID	A2MG MOUSE			
AC	P28665;			
DT	01-DEC-1992 (REL. 24, CREATED)			
DT	01-NOV-1995 (REL. 32, LAST SEQUENCE UPDATE)			
DT	01-NOV-1995 (REL. 32, LAST ANNOTATION UPDATE)			
DE	MURINOGLOBULIN 1 PRECURSOR (MUG1).			
GN	MUC1 OR MUG-1.			
OS	MUS MUSCULUS (MOUSE).			
OC	EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;			
OC	EUTHERIA; RODENTIA.			
RN	[1]			
RP	SEQUENCE FROM N.A., AND SEQUENCE OF 28-57.			
RC	TISSUE=LIVER;			
RX	MEDLINE; 91358495.			
RA	OVERBERGH L., TORREKENS S., VAN LEUVEN F., VAN DEN BERGHE H.;			
RL	J. BIOL. CHEM. 266:16903-16910(1991).			
CC	-I- FUNCTION: A PROTEINASE ACTIVATES THE INHIBITOR BY SPECIFIC			
CC	PROTEOLYSIS IN THE BAIT REGION, WHICH, BY AN UNKNOWN MECHANISM			
CC	LEADS TO REACTION AT THE CYSTEINYL-GLUTAMYL INTERNAL THIOLESTER			
CC	SITE AND TO A CONFORMATIONAL CHANGE, WHEREBY THE PROTEINASE IS			
CC	TRAPPED AND/OR COVALENTLY BOUND TO THE INHIBITOR. WHILE IN THE			
CC	TETRAMERIC PROTEINASE INHIBITORS STERIC INHIBITION IS SUFFICIENTLY			
CC	STRONG, MONOMERIC FORMS NEED A COVALENT LINKAGE BETWEEN THE			
CC	ACTIVATED GLUTAMYL RESIDUE OF THE ORIGINAL THIOLESTER AND A			
CC	TERMINAL AMINO GROUP OF A LYSINE OR ANOTHER NUCLEOPHILIC GROUP ON			
CC	THE PROTEINASE, FOR INHIBITION TO BE EFFECTIVE.			
CC	-I- TISSUE SPECIFICITY: PLASMA.			
CC	-I- SUBUNIT: MONOMER.			
CC	-I- SIMILARITY: TO OTHER PROTEINS OF THE ALPHA-MACROGLOBULIN FAMILY,			
CC	INCLUDING COMPLEMENT COMPONENTS C3, C4, AND C5.			
DR	EMBL; M65736; G199891; -.			
DR	PIR; A41185; A41185.			
DR	PROSITE; PS00477; ALPHA_2 MACROGLOBULIN.			
KW	SERINE PROTEASE INHIBITOR; GLYCOPROTEIN; PLASMA; BAIT REGION; SIGNAL;			
KW	MULTIGENE FAMILY.			
FT	SIGNAL 1 27			
FT	CHAIN 28 1476			
FT	DOMAIN 677 734			
FT	BAIT REGION (APPROXIMATELY)			
FT	MURINOGLOBULIN 1.			

FT	DISULFID	48	86	BY SIMILARITY.
FT	DISULFID	251	276	BY SIMILARITY.
FT	DISULFID	269	288	BY SIMILARITY.
FT	DISULFID	461	555	BY SIMILARITY.
FT	DISULFID	587	773	BY SIMILARITY.
FT	DISULFID	634	680	BY SIMILARITY.
FT	DISULFID	849	885	BY SIMILARITY.
FT	DISULFID	923	1323	BY SIMILARITY.
FT	DISULFID	1081	1129	BY SIMILARITY.
FT	DISULFID	1354	1469	BY SIMILARITY.
FT	THIOLEST	974	977	BY SIMILARITY.
FT	CARBOHYD	55	55	POTENTIAL.
FT	CARBOHYD	294	294	POTENTIAL.
FT	CARBOHYD	313	313	POTENTIAL.
FT	CARBOHYD	500	500	POTENTIAL.
FT	CARBOHYD	749	749	POTENTIAL.
FT	CARBOHYD	776	776	POTENTIAL.
FT	CARBOHYD	871	871	POTENTIAL.
FT	CARBOHYD	993	993	POTENTIAL.
FT	CARBOHYD	1142	1142	POTENTIAL.
FT	CARBOHYD	1180	1180	POTENTIAL.
FT	CARBOHYD	1426	1426	POTENTIAL.
SQ	SEQUENCE	1476	165139 MW; 7E3EAF00 CRC32;	

[illegible]

RESULT	15
ID	A2MG MOUSE
PRT;	1476 AA.
STANDARD;	
AC	P28665;
DT	01-DEC-1992 (REL. 24, CREATED)
DT	01-NOV-1995 (REL. 32, LAST SEQUENCE UPDATE)
DT	01-NOV-1995 (REL. 32, LAST ANNOTATION UPDATE)
DE	MURINOGLUBULIN 1 PRECURSOR (MUGI).
GN	MUC1 OR MUG-1.
OS	MUS MUSCULUS (MOUSE).
OC	EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
CC	EUTHERIA; RODENTIA.

RP	SEQUENCE FROM N.A., AND SEQUENCE OF 28-57.
RC	TISSUE=LIVER;
RX	MEDLINE; 91358495.
RA	OVERBERGH L., TORREKENS S., VAN LEUVEN F., VAN DEN BERGHE H.;
RL	J. BIOL. CHEM. 266:16903-16910(1991).
CC	-I- FUNCTION: A PROTEINASE ACTIVATES THE INHIBITOR BY SPECIFIC
CC	PROTEOLYSIS IN THE BAIT REGION, WHICH, BY AN UNKNOWN MECHANISM
CC	LEADS TO REACTION AT THE CYSTEINYL-GLUTAMYL INTERNAL THIOL ESTER
CC	SITE AND TO A CONFORMATIONAL CHANGE, WHEREBY THE PROTEINASE IS
CC	TRAPPED AND/OR COVALENTLY BOUND TO THE INHIBITOR. WHILE IN THE
CC	TETRAMERIC PROTEINASE INHIBITORS STERIC INHIBITION IS SUFFICIENTLY
CC	STRONG, MONOMERIC FORMS NEED A COVALENT LINKAGE BETWEEN THE
CC	ACTIVATED GLUTAMYL RESIDUE OF THE ORIGINAL THIOL ESTER AND A
CC	TERMINAL AMINO GROUP OF A LYSINE OR ANOTHER NUCLEOPHILIC GROUP ON
CC	THE PROTEINASE, FOR INHIBITION TO BE EFFECTIVE.
CC	-I- TISSUE SPECIFICITY: PLASMA.
CC	-I- SUBUNIT: MONOMER.
CC	-I- SIMILARITY: TO OTHER PROTEINS OF THE ALPHA-MACROGLOBULIN FAMILY,
CC	INCLUDING COMPLEMENT COMPONENTS C3, C4, AND C5.
DR	EMBL; M65736; G199891; -.
DR	PIR; A41185; A41185.
DR	PROSITE; PS00477; ALPHA_2_MACROGLOBULIN.
KW	SERINE PROTEASE INHIBITOR; GLYCOPROTEIN; PLASMA; BAIT REGION; SIGNAL;
KW	MULTIGENE FAMILY.
ET	SIGNAL 1 27
FT	CHAIN 28 1476
FT	DOWAIN 677 734
FT	BAIT REGION (APPROXIMATELY)











Qy	478	VGEHLNI-I-VTPKSPYIDKITHYNYLILSKGKIHFGRTRKFSDSAYOSINIPVTONWV	535
Db	518	psfrfayvyqy-gnn---eivadsvwvdvdkctcmgtllvvkaesrddrihqpaamkikle	574
Qy	536	PSRLLLVYIVTGEQTAELVSDSVLWNLIEEKGNQLOVH-LSPDADAY-SPQQTVSLNMA	593
Db	575	gdbqavglvavdkayvlnkykieqakiwdtiekedfgctagsgnmlgvfedaglal	634
Qy	594	TGMSWVALAANDVSAVYGVQRGAKEPLERVQFLEKSDJGCGAGGLNNAVYFHLGLTF	653
Db	635	tstetlnktksaakcpqpnrrrsrvllldsaskaaqfgdgglrkcecdgmhndpmg	694
Qy	654	LTWANADSQNDPECKE-IILRPRT---LQKITEEIAKYKHSVKKCYDGAQVWNDE	709
Db	695	yteckrakyiqedackaaflecchbykigidenqteselflaredfedelfgdnniier	754
Qy	710	-TCEQRAARISLPGRCIKAFTECCVWASQLRA-NISHKOMLGRHLMKLTPLVPVKPEI	766
Db	755	sdfeeswltteeltgepompngissitvpfyrlrdsttwellavlgleptkgicvaepyei	814
Qy	767	SYTPESWLEVH-LV-PR-R---RQLQFALPDSLTUWQIGIGISNT-GICVATVRA	817
Db	815	twmkdfidrlrplysvwnqeivrallynadedi-y-vr---velinyafcaesateg	869
Qy	818	KVFKDVFLENNIFYSVRGEQIQIKGTVYVRYRTSGMQFCVKMSAVEGICTSESPVDHQC	877
Db	870	qrv-r-qsfpkalsaravfivplegldhveiasvrgelaesdgvrkkllvvpeger	927
Qy	878	TKSKCVQKQVEGSSHLVFTVPLGLELNNFSLFW--FGKEILWKTLLVWPEGVK	935
Db	928	knivtielddpsvkvy-gtqeltviankid-kvdpdtevetrisvlgdpvaqiensid	985
Qy	936	RESYSGVTLDP--RGYGTISRKEFPRIPLDLPVKTEIKRIILSVKGLVGEILSAVLIS	993
Db	986	gsklnhliitpsgcgegmmitcpsviaty-yldatqwenlgvdrtte-a-ikqimt-g	1041
Qy	994	QEGINILTHLPKGSAAELMSVWP-VFYVYHLETGNHNNIFHSDPLIEKQIKKKIKG	1052
Db	1042	yacgmvykkadhyaafatnraseswltayvvkvyllanasmwkdshheiligvkvilnr	1101
Qy	1053	MLSIMSRYNADYSYVWKGSGASMTLAFALRVLIGQWVKYV-EQNQNSICNSLLWVNY	1111
Db	1102	qcpdgvfkenapvihgemlgtkgaep-eaal---tafiavtalleersvceqinildsei	1156
Qy	1112	QLDNGSFKENSQ-YQP IKLOGLTLPVARENSLYLTAFTVIGIRKAFDGC-PLWKI-DTAL	1168
Db	1159	nkadylllkvkeqlqrpvtaltayalaaadrlndd-r-v-v--lm-aa-stgrn--r-w	1207
Qy	1169	IKADNFFLENTLPAQSTFTTLAISAYALSIGKTHPQFRISVSAKREALWGNPPIVRFW	1228
Db	1208	e--ey-natthn-----iegtesyalllkmkkfaevgppvrvwllidkyvgtygqtgc	1257
Qy	1229	KONLQHKDSSVPNTGTARMVETTYATALLTNLKADINYYNVPVWKWLEEQRYGGFYSTQ	1288
Db	1258	atvmvfqalaeyeiqmphqdnlndisiklpetrevperysindrnvqartvecklneof	1317
Qy	1289	DTINAIEGLTEYSL-LVQKRLRLSMDIDVSYKHKGALHNYKMTDKNFL-GRPVEVLLNDL	1346
Db	1318	twase-gdgkatmtiltvnaqlredanvcnkhldvsvenvnlkqkggkaalrki	1376
Qy	1347	IUSTGFGSGLATVHVHTVWH-KYSTSEEVCS-FYKIDQDQIEASHYRVCNGSDYKRIVA	1404
Db	1377	ctry-lq--evds-tm-tiidiemltgfddaedklrknvgdtryiskfeidnmnackot	1431

[illegible]



FT /label= Sig\_peptide  
 FT Protein 23..1667  
 FT /note= "C3 beta chain"  
 FT Peptide 668..671  
 FT /note= "amino acids 668-671 are removed when the  
 FT precursor is cleaved into the alpha and  
 FT beta chains"  
 FT Protein 672..1663  
 FT /note= "C3 alpha chain"  
 FN #09607738-A2.  
 PD 14-MAR-1996.  
 PR 08-SEP-1995; G02121.  
 PR 08-SEP-1994; GB-018147.  
 PR 04-MAY-1995; GB-009102.  
 PA (IMUT-) IMUTRAN LTD.  
 PI Farries TC, Harrison RA;  
 DI WPI; 96-171613/17.  
 DR N-PSDB; T17738.  
 DR Mutant complement pathway protein forming stable C3 convertase -  
 PT for generalised complement depletion or localised complement  
 PT activation  
 PS Disclosure; Fig 1; 81pp; English.  
 CC Human C3 protein (R94028) was produced by expression of a cDNA  
 CC sequence (T17738) isolated from a human liver cDNA library.  
 CC C3 is a complement pathway protein that is acceptable to cleavage  
 CC by Factor I and is also susceptible to the inhibitory action  
 CC of Factor H. Mutants of C3 (R94029 and R94030) have been  
 CC produced by site-directed mutagenesis. These mutants can be  
 CC used to super-active the complement system, or to induce  
 CC localised super-activation at a specific target to increase  
 CC the target's sensitivity to complement-mediated destruction.  
 SQ Sequence 1663 AA;

Query Match 19.4%; Score 2339; DB 16; Length 1663;  
 Best local Similarity 28.3%; Pred. No. 1.02e-194;  
 Matches 488; Conservative 458; Mismatches 661; Indels 115; Gaps 90;

Db 8 sll-ll-llthlpalag-spmsiitpnllrseetmyleahdagqdvptvtvndfpg 64  
 Qy 2 GLLGILCFLIFLKGWQEQTYWISAPKIFRVGASENIVIQVGYTEAFATISIKSYD 61  
 Db 65 kkvlsesektvlpachnmgvntftipanrefkeekgrnkfvvtqatftgqve-kvvlv 123  
 Qy 62 KKFYSYSGHVHLSSENKFTNSAIIITQIP-KQLPGQNPVSYVYLEVV-SKHFPSKRWPI 119  
 Db 124 slsgyflqtdktytpqstvlrftvnhklllvgrtvmvniempgipvkqdslesq 183  
 Qy 120 TYDNGELEIHTDKPVYTPDQSKVRVYSINDDLKPAKRETVLFTIDPESEV--DWVEEI 177  
 Db 184 nqlvlples-wdipeivmngkwirayenspqqvfstefevkvlpsfevipeptekf 242  
 Qy 178 DHGIISLPPDFKIPSNRYGMMWIKAKYREDFSTGTAYFEVKEYVLPHFSVTEP-E-Y 235  
 Db 243 yyl-yne-kglevtitarflygkkv-egtatvfigi-qd--geqrialspeelkripledg 296  
 Qy 236 NFICYKNFQNFETIKARYFYFNWVTEADVITFGIREDIADKQKEMQMTAMONTMLNG 295  
 Db 297 sgevlrsrkvllqgvmpnaedilvgklylsatvilhsesdmvqaersipivtpeyqih 356  
 Qy 296 IAQVTFDSETAVKELSYSLLEDLNKKYLYIAVTVIESGTGFSSEAEIPGKYKVLSPYKLN 355  
 Db 337 ftktpkykpmqfdlmvfnvp-d---gs-pay-rvp-vav-q--ge-dtvqslq-qd 404  
 Qy 356 LVATPLFLKPGIDPPKQVQKSDLDLUGGVPIVLAQITDINQETSDLDPSKSVTRVDD 415

Db 405 qvaklsinthpsqkplsvtrtkkqelseaeatrtmqalpystvtqsnnylnhslvrte 464  
 Qy 416 GVASFVINLPISGVTVLEFNVKTDAPDIPFENQAREGYRAIAYSSLSQSYLIDWTDNIKA 475  
 Db 465 lrpgelinvfillrmdraheakirytyliannkgrllkagrqvrepqgdlvvlplaitd 524  
 Qy 476 LLVGEHLAI-IVTPKS-PYIDKITHYILILSKGKIHFCTREKESDASYSQISINPVTON 533  
 Db 525 fipefrlvayvtlligasqrevadsvwdvdkdcvqslvksqgsedrtprvpgqmtlk 584  
 Qy 534 MWPSSRLVYIVTGEQTA-ELVSDSVMLNTEEXCGNQLQVHLSPDADAYS-PCQTVSLN 591  
 Db 585 iegdgarvllvavdgfvlnknkltqskiwdvvekadictpggkdyagvdsdagl 644  
 Qy 592 MATCMDSWVALAAVDNAVYGVQRCAGKPLERVFQFLEKSDLCGGAGGLNNANVPHLAGL 651  
 Db 645 tftsssgqgtatraelqcpqpaaarrrrsvqltekrmdkvq-kypkel-rkccedgmtenp 702  
 Qy 652 TFLTNANADDSQENDEPCKE-ILPRRT--L-QKKIEEIAAKYKHSVVKCCYDGCACVNN 707  
 Db 703 mrfscqrtrfislgeackvflcdcnymitelrrqharashlglaranidediaeeniv 762  
 Qy 708 DE-TCQRAARISIGPRCTKAFTECCVWASQIRANISH-KDMQLGRLLHK-TLLPVSKE 764  
 Db 763 srsefpewlnvnedikeppknigistklmnlfldsittweilavmsdkkgicvadpfe 822  
 Qy 765 IRSYFESWMEVH-L-VPRRK---QL-QFALPDSLTTWEIQGIGISNT-GICVADTVK 816  
 Db 823 vtvmqdfidrlpswvvrneqveiravlynrgnqgelkvrvellhnpafcsatlattkrh 882  
 Qy 817 AKVKDFVLEWNTIPYSVVRGEQIQKCTVYNYRTS-QMQFCVKMSAVEGICITESPVYDH 875  
 Db 883 qqtit---ippk--sals--vpyvivplktglqeavkaavyhhfisdgvrkslkvypge 935  
 Qy 876 QGTSKSKVRQKVEGSSSHLVTFVLPLEIGHNINFSLETW--FGKEILVKTILRVVPEG 933  
 Db 936 irmnktavrtldperl-gregvqkedip-padisdqvdtesetrlil-lqgtpvagmte 992  
 Qy 934 VKRESYGV-TLDPRGITISRKE-FPYRIPL-DLVPKTEI-KRILSVKGLLVGEILS 989  
 Db 993 davdaerlkhliutpgcgeqnmigmtpctviahvyldeqewkf-q--lekrgqaleli 1049  
 Qy 990 AVLSQGINILTHLPKGSAAELMSVVPVYVYHYLETGNHWNIFHSDPLIEKQKLKKKL 1049  
 Db 1050 kkgvtqqlafrcpsaafvkrapatwltayvykvsfslavnlaidsvlcvavkwlll 1109  
 Qy 1050 KGLMSINSYENADYSYVWKGSSASTMTLTAFLRVLGQVNYEQNQNSICNSLJLWVE 1109  
 Db 1110 ekqtkpvgfedapvihgemiqgllr-nnnnekdmaltafvlsalgeakdiceevnslpgs 1168  
 Qy 1110 NYQDNGSFKNESQYQPIKIAQGTLPVEARENSLYLTAFTVIGIRKAFDICP-LVK-IDTA 1167  
 Db 1169 itkagdfiaanymlqrsyvtvaiaqvala--qmgr--lkg--pllnk-flttakdk-nr- 1219  
 Qy 1168 LTKADNFLENTLPAQSTFTLTAISVALSLGSKTHPQFRSVSALKREALVKGNPPYRF 1227  
 Db 1220 w-ed-pgkq--l-yn-----veatsyallallqlkdfdvppvvrwlnesqrvyggvgst 1269  
 Qy 1228 WKONLQHKDSSVPNTGTARWVETTAVALLTSLNLKIDINYVNPVIKWLEEQRYGCGFYST 1287  
 Db 1270 qatfmvqalaqkdpdthqelnldvsllqpsrskithrihwesasllreetkeneg 1329  
 Qy 1288 QDTINAEGLTEISL-LVKQLRLSMDIDVSYKHKGALHNYKWT-DKNFLGRPVEVLNDD 1345





Qy 240 YGNFK-NFEITIKARYFNKVVTEADVYITFGIRELDKQKEMQATQNTMLINGIAQ 298  
Db 292 atldrb-frs-rfpnlmqlvghtlvaavtmtesgdmvvtqsgihivaspyqlhftk 349  
Qy 299 VTFDSEATKELSYSLIEDINNYLYIAVTVIESTGSEAEIPGKGVKLVKLVIA 358  
Db 350 tpykfpmpyeltvvtvnp-d--gs-paa-hvp-v-v-ea--fh-smgtt--ledgta 396  
Qy 359 TPLFKPGIPYPIKQVQKDSLDLQVGPVVLNAQTIIDVNOETSDLDPSKSVTVDDGVA 418  
Db 397 klilnplnagslpitvrthngldprerqatksmtaiaqvtqgsgnylhvaiteiekp 456  
Qy 419 SFVILPESGVTVLEFNKVTOPALPEENQAREGYRAIAYSLSQSYLYIDWTDNHRKALV 478  
Db 457 gdnlpvfnvknanslkqikyftylilnkqikfkvgqrrdqgnlvmtlmhltpdilip 516  
Qy 479 GERINI-I-VTPKSPYIDKITHNYLILSKGKIHFETREKFSDAYSQISINIPVQNWVP 536  
Db 517 srfvayqv-gnn--eivadsvwvdktdcmgtl-v-vkgdnliimgpaamkiklegdp 571  
Qy 537 SRRLLVYVYTGEOIAELVSDSVINIEKCGNQLQVHLSPDADAISPCQTVSLNWATCH 596  
Db 572 gatvlgavdkavylndykieqakidwtiekedfgctagsgmnlgvfedaqlatts 631  
Qy 597 DSWALANVSAYGVQBAKKELEVRQFLEKSDGCGGAGGLNANVFLHGLIFLTLN 656  
Db 632 tnlnkrgsaakpnparrirrsvllldenaakaefqddlrkccedvmhnpngytc 691  
Qy 657 ANADDSQENDEPCKE-ILRPRT---LQKTEEIAAKYKHSWKCCYDGCAGVNDE-TC 711  
Db 692 ekrazyiqedackaafleccrykyvndenqreselflarddnedgdiadsidf 751  
Qy 712 EQRAARISLGRICAKFTECCVVASQLRA-NISHKDMQLGRL-HMKTLLPVSKPEIRSYF 769  
Db 752 pkewlwtldlteepnsggisaktsmfylrdsittwv-lavsfptkicvaepyeirvm 811  
Qy 770 PESWLEW-HLV--PR-R---KQQLFALPDLSTWETQIGISINT-GICVADTVKAKVF 820  
Db 812 kvffidlmpysvknegveirailhnyvmedl-y-vr---velly---npafcsastkq 863  
Qy 821 KDVLENNIPYSVWRGEQIQKGTVTYNTYRSGMQFCVMSAVEGICTSESPIVDHQGTKS 880  
Db 864 qryrqfpikaasrtavpvlvpleglhdveikaavgealw-edgv-rkklkvpeqvq 921  
Qy 881 SKCVRQ-KVEGSSHLVFTPLPLEIHLNINF--SL-ET-WFGKEILVKTIRVWPEGVK 935  
Db 922 ksvitvklprak-gvogtqlvrikardldrvpdeietkiiigdpdvaqiensidg 980  
Qy 936 RESYSGVTLDPRICTYTSRRKE-PPYRIPLDLVLPKTEIKRILSVKGLVGEILSAVLQS 994  
Db 981 sklnhlitpscgqgmirmaaapviaty-yldtqwetlginrte-a-vnqivt-gy 1036  
Qy 995 EGINILTHLPKSAEAEMLSW-PVYFVHYLETGNHWNIFHSDPLIEKQKLKKLKEGM 1053  
Db 1037 aqmqvkkadhysaaftrasswltavvkvfamaakmvgihsellicgqvrvlllnrq 1096  
Qy 1054 LSIHSTRNADYSYVWKGSGSALTWLFAURLVQGNKYVEQ-NQNSICNSLMLWENVYQ 1112  
Db 1097 qpdgaftenapvlsgtmagg--iqgaeevyltafilvalleakticndymldessikk 1154  
Qy 1113 LDNGSFENSOYQIKLAGTLPVEARENSLYLTAFTVIGIRKAFDTCF-LVK-IDTALIK 1170  
Db 1155 atnyllkkyelqrpyttaltayalaadqldnd-r-v--lm-aa-stgrd--h-w-e 1202

Qy 1171 ADFNLTLPAQSTFTLAIASVALSIGDKTHPQFSRVSALAKREALVKGNPPIYREWKD 1230  
Db 1203 --ey-na--ht---hniegtcyalllkmkfdtqgpiwrltdqmfyetygqtat 1253  
Qy 1231 NIQHKSSVPNTGTARWETAYALLTSLNKDINYNPVIKWLSEQRVGGGFYSTODT 1290  
Db 1254 vmafqalaeeygmphthkldnlditelpdevprryinyenallartvetklnqditv 1313  
Qy 1291 INAIEGLEFTEYSL-LVKQLRLSMDIDVSYKHKALHNYKWT-DKQFLGRPVEVLNDDLLIV 1348  
Db 1314 tas-gdakatiltfynaqlqekavcnkfhlnvsvenihlna-mg-akgalmkiki-ct 1369  
Qy 1349 STGFGSLGTLVHTVTVVH-KTSTSEEVCS-FYLIKIDTQDIEASHYRGYNSDKRIVACA 1406  
Db 1370 ry-lg--evds-tm-tiidismltqfipdaedlrrlekvgdryisryevdhnmaqvavi 1424  
Qy 1407 SYKPSREESSGSHAVMDISLPTGISANEEDIKALIVEGVQDLFTDYQI-KD-GH---VI 1461  
Db 1425 iylnkvshsdeclhfklnhfevgfigpagsvkvysynldekctkfyhpdkggtllnki 1484  
Qy 1462 LQJNSIPSSDFLCVREIRIFELFEVGLSPATFTVYEHPRDKQCTMFYSTSN- IKI-QKV 1519  
Db 1485 cignvrcagatcsslnhq-e-rldvplqiekacetnvdvyvktklrlieeqdndiyvm 1542  
Qy 1520 CEGACKCEADCGQMQGELOLTISAEKTKACKETAIYAKVSIITSITVENFVKYKA 1579  
Db 1543 dvlevikgtdknprakthqvisqrkcealnkvmdvdyliwgsrdliptk-d-kisyi 1600  
Qy 1580 TLLDIYTKGEAVAEKQSEITFKVKVCTNA-ELVKGQYLLMG-K-EALQIKYNSFRYI 1636  
Db 1601 it-kn-twierwhbedeqe-eef-qklcddfaq 1630  
Qy 1637 YPLDSLUTWYMPRDTTSSQQAFLANL-DEFAE 1669  
RESULT 7  
ID R63224 standard; Protein; 1333 AA.  
AC R63224;  
DT 05-JUL-1995 (first entry)  
DE Cobra partial CVF2.  
KW Cobra; C3; third component of complement; human; mouse; rat;  
KW X. laevis; pre-pro molecule; beta chain; alpha chain; codon usage;  
KW immune response; host defence; tumour.  
OS Naja naja.  
FH Key Location/Qualifiers  
FT Region 1..332  
FT /note= "Partial alpha chain"  
FT Region 416..715  
FT /note= "Gamma chain"  
FT Region 947..1333  
FT /note= "Beta chain"  
PN W09423024-A.  
PD 13-OCT-1994.  
PF 07-APR-1994; U03441.  
PR 07-APR-1993; US-043747.  
PA (GEOU ) UNIV GEORGETOWN.  
PI Bredehorst R, Fritzinger DC, Vogel C;  
DR N-PSDB; Q77791.  
PT DNA encoding cobra C3, CVF 1 and CVF 2 - which are used in the  
PT treatment of cancer  
PS Claim 5; Fig 2M-2V; 155pp; English.  
CC This sequence represents the C-terminal portion of cobra venom factor



977 LSXVGLVGEJLSAVLSQEGINLILHFPAGSAEALMSVV-PVFIVRHLEIGNHNIIFH 1033

FI Esper B, Latis 3-0;  
DR WPI; 91-102075/14.







PD 15-JUN-1995.  
PF 16-NOV-1994; 1B0359.  
PPR 06-DEC-1993; US-162591.  
PR (CIBA ) CIBA GEIGY AG.  
PPI Boyar WC. Galakatos NG, Peppard JV, Van Oostrum J;  
DR WFI; 95-224319/29.  
DPI N-PSDB; 092518.  
DPT C5a receptor antagonists having no agonist activity - are used in  
DPT compans. to treat C5a-mediated diseases and inflammatory conditions  
DPT Disclosure; Page 36-37; 65pp; English.  
CC C5a encoded by a synthetic gene is given in R75497. Analogs of C5a,  
CC obtd. by mutagenesis of the C-terminal region, exhibit excellent  
CC antagonistic properties and substantially no agonist activity.  
CC Sequence 74 AA;  
ISO

[illegible]

Search completed: Wed Jan 28 12:13:50 1998  
Job time : 66 secs.

[illegible]

RESULT	13	
AD	R21775 standard; Protein; 344 AA.	
AC	R21775;	
DD	25-JUN-1992 (first entry)	
DT	Phospholipase A2 inhibitory protein.	
DE	Inflamed tissue; antihistamine; allergic reactions.	
OW	Rattus rattus.	
OS	W09202619-A.	
PN	20-FEB-1992.	
PD	03-AUG-1991; J01040.	
PF	03-AUG-1990; JP-205164.	
PR	19-DEC-1990; JP-411594.	
PRR	(TEIJ ) TEIJIN KK.	
PA	Suwa Y, Imaizumi A, Okada M, Suzuki Y, Kudo I, Inoue K;	
PI	Azumac, Murakami M;	
PT	WPI; 92-080073/10.	
OR	N-PSDB; Q21650.	
DB	Plasmid encoding phospholipase A2 inhibitory protein - derived	
PTT	from human or rat inflamed region, used as antihistamine agents	
TT	for treating allergic disorders	
PTT	Disclosure; Page 31; 52pp; Japanese.	
PS	A rat liver cDNA lambda gt11 library obtd. from inflamed tissue	
CC	was screened using mouse C3cDNA-derived plasmid pFC4/5.4 (cleaved	
CC	with HindIII/StuI) as probe. The rat C3cDNA clone pTC3/11 obtd.	
CC	was cleaved with EcoRI into 2.0 kb and 0.1 kb fragments which	
CC	were combined with vector pTV119N to give pPTC3/2.1. The plasmid	
CC	was used to transform E. coli and the transformant found to express	
CC	a phospholipase inhibitory protein of Mw. 39 kD (SEQ ID NO 1). The	
CC	protein can be used as an anti-histamine agent which can be used	
CC	to treat allergic reactions.	
CC	See also R21776.	
CC	Sequence 344 AA;	
CC	30	

[illegible]

	1145	TAFTVIGIRKAFDIP-LVK-IDTALIKADNFLELNTLPASQSTFTTLAISAYALSUDKTH	1202
Ddb	245	epy--lktflint-a-k-dpn--r-weepgqg----l-yn-----veatsyallalililk	285
	:	:   :   :   :   :   :   :	:   :   :   :
Qoy	1203	PQRSTVSALKREALVKNPPIYRFWKONLQHDKSSVPNTGTRARWETATAYALLTSNIK	1262
Ddb	286	dfdsvppvrvlnearyyvgggystgatfmvfgalaagcqtvdphdkdlmndval	339
	:	:   :   :   :   :   :   :	:   :   :   :
Qoy	1263	DINYNPVIKWLEEQRYGGGFYSTQDTINAIEGLTEYSLLVKQLR-LSWDIDV	1315
	:	:   :   :   :   :   :   :	:   :   :   :
	RESULT	14	
ID	R10899	standard; Protein; 344 AA.	
AC	R10899;		
DT	07-MAY-1991	(first entry)	
DE	Rat phospholipase A2 inhibitor protein.		
KW	Inflammatory disease.		
OS	Rattus rattus.		
PN	MO9101999-A.		
PD	21-FEB-1991.		
PF	03-AUG-1990;	J00996.	
PR	03-AUG-1989;	JP-200246.	
PPR	05-APR-1990;	JP-089085.	
PPA	(TEIJ ) TEIJJN KK.		
PPI	Sawa Y, Imaizumi A, Okada M, Kudo I, Inone K, Suzuki Y;		
DR	WP1; 91-073488/10.		
DR	N-P5DB; Q10773.		
DR	Phospholipase A2 inhibitor protein - obtd. by enzymic treatment		
RPT	of mammalian serum.		
CC	Claim 1; Fig 1; 68pp; English.		
CCC	The protein, obtd. from rat serum or expressed from the gene, is		
CCC	used in the treatment and diagnosis of inflammatory diseases.		
CCC	See also R10900.		
CSQ	Sequence	344 AA;	

		Query Match	3.6%	Score 436;	DB 3;	Length 344;
		Best Local Similarity	28.2%;	Pred. No.	4.57e-25;	
		Matches	100;	Conservative	99;	Mismatches 127; Indels 28; Gaps 17;
bB	bb	10	dqvdptdsetril-lqgtpvgaadeavdglerkhilvtpgcgcgmigmtptviavhy	68		
			:     :  :::	:   :   :	:     :   :	
yY	yy	966	DLVPTKTL-KRILSVKGILLVGEILLSAVLSQEGINITHPLPGSSEAELMSVVVFYFHY	1024		
bB	bb	69	ldqtceqekf-g-lekrqealellikkgtqqqlafkqpessavaafnrrpstwtlctayvk	125		
			: ::   :   :   :	:   :   :   :   :	:	: :   :     :   :
yY	yy	1025	LETGNHNIHFSDPLEIKOKLIKKKLKEGMLSIMSYRNADYSVWKGGSASTMTAFALR	1084		
bB	bb	126	vfslaanliaidsqvlgacvwklliiekqpdqgfcdpqgvihqmieggr-ftnteadvsrl	184		
			: : : :   :   :   :	:   :   :   :	: :   :   :	
yY	yy	1085	VLGOWNKVEQNONSICNSLLWVENYQLDNGSFKENSOYPQIKLAGTLPVEARENSLYL	1144		
bB	bb	185	t:fvlliaigeardicedqvnslpsgekageyleasylnlqrpytvaigaalamlmkle	244		
			:     :   :   :	:   :   :   :	:   :   :   :	
yY	yy	1145	TAFTVIGIRKAEDICP-LVK-IDTALIKADFLENTLPASGTFTLAISAYALSIGDKTH	1202		
bB	bb	245	epy--ltckflint-a-k-drn--r-weepggq---lyn-----veateysalllllk	285		
			:   :   :   :	:   :   :   :	:	:   :   :   :
yY	yy	1203	POFRSIVSLAKREALVLKGNPPYPRFWKNOLQHDKSSVPHTCTARMMETTAYALLTSNLK	1262		
bB	bb	286	fdsdvppvvrvineqryvggygstqatfmvfqaalgagtcdvpdhkdmlmvsrl	339		
			:   :   :   :   :   :	:   :   :   :   :	:   :   :   :	
yY	yy	1263	DINYNPVIKLLEEQRGGYGSTQTDTINAIEGLTEYSLLVKKQLA-LSMDDIDV	1315		